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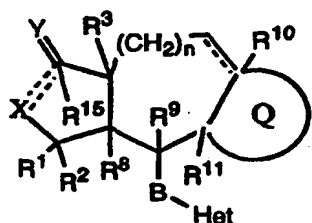
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<p>(51) International Patent Classification⁶ : C07D 405/06, 405/14, 409/14, 405/12, 417/06, 401/06, 491/04, 471/04, 493/10, A61K 31/44, 31/47 // (C07D 491/04, 317:00, 221:00) (C07D 471/04, 221:00)</p>	<p>A1</p>	<p>(11) International Publication Number: WO 99/26943 (43) International Publication Date: 3 June 1999 (03.06.99)</p>
<p>(21) International Application Number: PCT/US98/24523 (22) International Filing Date: 23 November 1998 (23.11.98) (30) Priority Data: 08/977,979 25 November 1997 (25.11.97) US (71) Applicant: SCHERING CORPORATION [US/US]; 2000 Galloping Hill Road, Kenilworth, NJ 07033-0530 (US). (72) Inventors: CHACKALAMANNIL, Samuel; 790 Stratford Road, East Brunswick, NJ 08816 (US). ASBEROM, Theodoros; 17 Stone Drive, West Orange, NJ 07052 (US). XIA, Yan; 137 Christie Street, Edison, NJ 08820 (US). DOLLER, Dario; 186 Filly Drive, North Wales, PA 19454 (US). CLASBY, Martin, C.; 231 Spruce Mill Lane, Scotch Plains, NJ 07076 (US). CZARNIECKI, Michael, F.; 30 Gildersleeve Place, Watchung, NJ 07060 (US). (74) Agents: MAGATTI, Anita, W. et al.; Schering-Plough Corporation; Patent Dept., K-6-1 1990, 2000 Galloping Hill Road, Kenilworth, NJ 07033-0530 (US).</p>		<p>(81) Designated States: AL, AM, AU, AZ, BA, BB, BG, BR, BY, CA, CN, CZ, EE, GD, GE, HR, HU, ID, IL, IS, JP, KG, KR, KZ, LC, LK, LR, LT, LV, MD, MG, MK, MN, MX, NO, NZ, PL, RO, RU, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UZ, VN, YU, ARIPO patent (GH, GM, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i></p>

(54) Title: **THROMBIN RECEPTOR ANTAGONISTS**



(1)



(A)



(B)

(57) Abstract

Heterocyclic-substituted tricyclics of formula (I) or a pharmaceutically acceptable salt thereof, wherein: the single dotted line represents an optional double bond; the double dotted line represents an optional single bond; n is 0-2; Q is optionally substituted cycloalkyl, heterocycloalkyl, aryl or heteroaryl; Het is an optionally substituted mono-, bi- or tricyclic heteroaromatic group; B is $-(CH_2)_{n3}-$, wherein n_3 is 0-5, $-CH_2-O-$, $-CH_2S-$, $-CH_2-NR^6-$, $-C(O)NR^6-$, $-NR^6C(O)-$, (a), optionally substituted alkenyl or optionally substituted alkynyl; X is $-O-$ or $-NR^6-$, when the double dotted line represents a single bond, or X is $-OH$ or $-NHR^{20}$ when the bond is absent; Y is $=O$, $=S$, (H, H), (H, OH) or (H, C_1-C_6 alkoxy) when the double dotted line represents a single bond, or when the bond is absent, Y is $=O$, (H, H), (H, OH), (H, SH) or (H, C_1-C_6 alkoxy); R^{15} is absent when the double dotted line represents a single bond and is H, $-NR^{18}R^{19}$, or $-OR^{17}$ when the bond is absent; or Y is (A) or (B) and R^{15} is H or C_1-C_6 alkyl; are disclosed, as well as pharmaceutical compositions containing them and a method of treating diseases associated with thrombosis, atherosclerosis, restenosis, hypertension, angina pectoris, arrhythmia, heart failure, and cancer by administering said compounds.

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THROMBIN RECEPTOR ANTAGONISTS

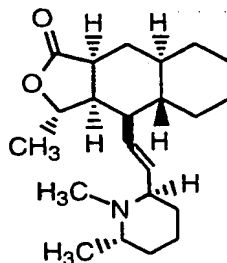
BACKGROUND OF THE INVENTION

The present invention relates to substituted tricyclic thrombin
10 receptor antagonists, pharmaceutical compositions containing them and
their use in the treatment of diseases associated with thrombosis,
atherosclerosis, restenosis, hypertension, angina pectoris, arrhythmia,
heart failure, cerebral ischemia and cancer.

Thrombin is known to have a variety of activities in different cell
15 types and thrombin receptors are known to be present in such cell types
as human platelets, vascular smooth muscle cells, endothelial cells and
fibroblasts. It is therefore expected that thrombin receptor antagonists
will be useful in the treatment of thrombotic, inflammatory,
atherosclerotic and fibroproliferative disorders, as well as other
20 disorders in which thrombin and its receptor play a pathological role.

Thrombin receptor antagonists peptides have been identified
based on structure-activity studies involving substitutions of amino
acids on thrombin receptors. In Bernatowicz et al, *J. Med. Chem.*, 39
(1996), p. 4879-4887, tetra- and pentapeptides are disclosed as being
25 potent thrombin receptor antagonists, for example N-trans-cinnamoyl-p-
fluoroPhe-p-guanidinoPhe-Leu-Arg-NH₂ and N-trans-cinnamoyl-p-
fluoroPhe-p-guanidinoPhe-Leu-Arg-Arg-NH₂. Peptide thrombin
receptor antagonists are also disclosed in WO 94/03479, published
February 17, 1994.

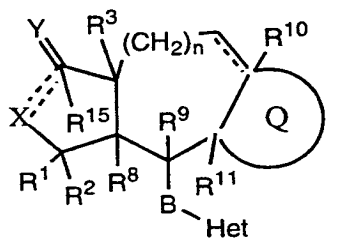
30 Himbacine, a piperidine alkaloid of the formula



has been identified as a muscarinic receptor antagonist. The total synthesis of (+)-himbacine is disclosed in Chackalamannil et al, J. Am. Chem. Soc., **118** (1996), p. 9812-9813.

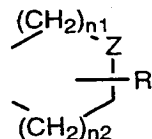
5 SUMMARY OF THE INVENTION

The present invention relates to thrombin receptor antagonists represented by the formula I



or a pharmaceutically acceptable salt thereof, wherein:

- 10 the single dotted line represents an optional double bond;
the double dotted line represents an optional single bond;
n is 0-2;
Q is



- 15 wherein n_1 and n_2 are independently 0-2; or when the double bond is not present, Q is also fused R-substituted aryl or R-substituted heteroaryl;

R is 1 to 3 substituents independently selected from the group consisting of H, C₁-C₆ alkyl, halogen, hydroxy, amino, (C₁-C₆)alkyl-amino, (C₁-C₆)dialkylamino, (C₁-C₆)alkoxy, -COR¹⁶, -COOR¹⁷, -SOR¹⁶, -SO₂R¹⁶, -NR¹⁶COR^{16a}, -NR¹⁶COOR^{16a}, -NR¹⁶CONR⁴R⁵, fluoro-(C₁-C₆)alkyl, difluoro(C₁-C₆)alkyl, trifluoro(C₁-C₆)alkyl, C₃-C₆ cycloalkyl, C₂-C₆ alkenyl, aryl(C₁-C₆)alkyl, aryl(C₂-C₆)alkenyl, heteroaryl(C₁-C₆)alkyl, heteroaryl(C₂-C₆)alkenyl, hydroxy(C₁-C₆)alkyl, amino(C₁-C₆)alkyl, aryl and thio(C₁-C₆)alkyl;

R¹ and R² are independently selected from the group consisting of H, C₁-C₆ alkyl, fluoro(C₁-C₆)alkyl, difluoro(C₁-C₆)alkyl, trifluoro-(C₁-C₆)alkyl, C₃-C₆ cycloalkyl, C₂-C₆ alkenyl, aryl(C₁-C₆)alkyl, aryl(C₂-C₆)alkenyl, heteroaryl(C₁-C₆)alkyl, heteroaryl(C₂-C₆)alkenyl, hydroxy-

(C₁-C₆)alkyl, amino(C₁-C₆)alkyl, aryl and thio(C₁-C₆)alkyl; or R¹ and R² together form a =O group;

R³ is H, hydroxy, C₁-C₆ alkoxy, -SOR¹⁶, -SO₂R¹⁷, -C(O)OR¹⁷, -C(O)NR¹⁸R¹⁹, C₁-C₆ alkyl, halogen, fluoro(C₁-C₆)alkyl, difluoro(C₁-C₆)alkyl, trifluoro(C₁-C₆)alkyl, C₃-C₆ cycloalkyl, C₂-C₆ alkenyl, aryl(C₁-C₆)alkyl, aryl(C₂-C₆)alkenyl, heteroaryl(C₁-C₆)alkyl, heteroaryl(C₂-C₆)alkenyl, hydroxy(C₁-C₆)alkyl, amino(C₁-C₆)alkyl, aryl or thio(C₁-C₆)alkyl;

Het is a mono-, bi- or tricyclic heteroaromatic group of 5 to 14 atoms comprised of 1 to 13 carbon atoms and 1 to 4 heteroatoms independently selected from the group consisting of N, O and S, wherein a ring nitrogen can form an N-oxide or a quaternary group with a C₁-C₄ alkyl group, wherein Het is attached to B by a carbon atom ring member, and wherein the Het group is substituted by 1 to 4 substituents, W, independently selected from the group consisting of H; C₁-C₆ alkyl; fluoro(C₁-C₆)alkyl; difluoro(C₁-C₆)alkyl; trifluoro-(C₁-C₆)-alkyl; C₃-C₆ cycloalkyl; heterocycloalkyl; heterocycloalkyl substituted by C₁-C₆ alkyl or C₂-C₆ alkenyl; C₂-C₆ alkenyl; R²¹-aryl(C₁-C₆)alkyl; R²¹-aryl-(C₂-C₆)-alkenyl; heteroaryl(C₁-C₆)alkyl; heteroaryl(C₂-C₆)-alkenyl; hydroxy(C₁-C₆)alkyl; dihydroxy(C₁-C₆)alkyl; amino(C₁-C₆)alkyl; (C₁-C₆)alkylamino-(C₁-C₆)alkyl; di-((C₁-C₆)alkyl)-amino(C₁-C₆)alkyl; thio(C₁-C₆)alkyl; C₁-C₆ alkoxy; C₂-C₆ alkenyloxy; halogen; -NR⁴R⁵; -CN; -OH; -COOR¹⁷; -COR¹⁶; -OSO₂CF₃; -CH₂OCH₂CF₃; (C₁-C₆)alkylthio; -C(O)NR⁴R⁵; -OCHR⁶-phenyl; phenoxy-(C₁-C₆)alkyl; -NHCOR¹⁶; -NHCO₂R¹⁶; biphenyl; -OC(R⁶)₂COOR⁷; -OC(R⁶)₂C(O)NR⁴R⁵; (C₁-C₆)alkoxy; C₁-C₆ alkoxy substituted by (C₁-C₆)alkyl, amino, -OH, COOR¹⁷, -NHCOOR¹⁷, -CONR⁴R⁵, aryl, aryl substituted by 1 to 3 substituents independently selected from the group consisting of halogen, -CF₃, C₁-C₆ alkyl, C₁-C₆ alkoxy and -COOR¹⁷, aryl wherein adjacent carbons form a ring with a methylenedioxy group, -C(O)NR⁴R⁵ and heteroaryl; R²¹-aryl; aryl wherein adjacent carbons form a ring with a methylenedioxy group; heteroaryl; heteroaryl substituted by 1 to 4 substituents selected from the group consisting of halogen, C₁-C₆ alkoxy, C₁-C₆ alkyl, (C₁-C₆)alkylamino, di-((C₁-C₆)alkyl)amino, -OCF₃, -NO₂, hydroxy(C₁-

C₆)alkyl, -CHO and phenyl; and heteroaryl wherein adjacent carbon atoms form a ring with a C₃-C₅ alkylene group or a methylenedioxy group;

- 5 R⁴ and R⁵ are independently selected from the group consisting of H, C₁-C₆ alkyl, phenyl, benzyl and C₃-C₆ cycloalkyl, or R⁴ and R⁵ together are -(CH₂)₄-, -(CH₂)₅- or -(CH₂)₂NR⁷-(CH₂)₂- and form a ring with the nitrogen to which they are attached;


R⁶ is independently selected from the group consisting of H, C₁-C₆ alkyl or phenyl;

- 10 R⁷ is H or (C₁-C₆)alkyl;

R⁸, R¹⁰ and R¹¹ are independently selected from the group consisting of R¹ and -OR¹, provided that when the optional double bond is present, R¹⁰ is absent, and when ring Q is aromatic, R¹⁰ and R¹¹ are absent;

- 15 R⁹ is H, OH, C₁-C₆ alkoxy, halogen or halo(C₁-C₆)alkyl;

B is -(CH₂)_{n3}-, -CH₂-O-, -CH₂S-, -CH₂-NR⁶-, -C(O)NR⁶-.

-NR⁶C(O)-, , cis or trans -(CH₂)_{n4}CR¹²=CR^{12a}(CH₂)_{n5} or -(CH₂)_{n4}C≡C(CH₂)_{n5}-, wherein n₃ is 0-5, n₄ and n₅ are independently 0-2, and R¹² and R^{12a} are independently selected from the group

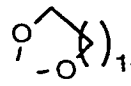
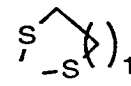
- 20 consisting of H, C₁-C₆ alkyl and halogen;

X is -O- or -NR⁶- when the double dotted line represents a single bond, or X is -OH or -NHR²⁰ when the bond is absent;

Y is =O, =S, (H, H), (H, OH) or (H, C₁-C₆ alkoxy) when the double dotted line represents a single bond, or when the bond is absent, Y is

- 25 =O, (H, H), (H, OH), (H, SH) or (H, C₁-C₆ alkoxy);

R¹⁵ is absent when the double dotted line represents a single bond and is H, C₁-C₆ alkyl, -NR¹⁸R¹⁹, or -OR¹⁷ when the bond is

absent; or Y is  ₁₋₂ or  ₁₋₂ and R¹⁵ is H or C₁-C₆ alkyl;

- 30 R¹⁶ and R^{16a} are independently selected from the group consisting of C₁-C₆ lower alkyl, phenyl or benzyl;

R¹⁷, R¹⁸ and R¹⁹ are independently selected from the group consisting of H, C₁-C₆ alkyl, phenyl, benzyl;

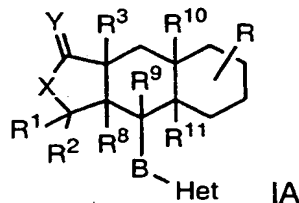
R²⁰ is H, C₁-C₆ alkyl, phenyl, benzyl, -C(O)R⁶ or -SO₂R⁶;

R²¹ is 1 to 3 substituents independently selected from the group consisting of -CF₃, -OCF₃, halogen, -NO₂, C₁-C₆ alkyl, C₁-C₆alkoxy, (C₁-C₆)alkylamino, di-((C₁-C₆)alkyl)amino, amino(C₁-C₆)alkyl, (C₁-C₆)-alkylamino(C₁-C₆)alkyl, di-((C₁-C₆)alkyl)-amino(C₁-C₆)alkyl, hydroxy-
 5 (C₁-C₆)alkyl, -COOR¹⁷, -COR¹⁷, -NHCOR¹⁶, -NHSO₂R¹⁶ and -NHSO₂CH₂CF₃;

Z is -CH₂-, -O-, -S(O)₀₋₂-, -NR²²-, -C(O)-, -C(=NOR¹⁷)- or -C(R¹³R¹⁴)-, wherein R¹³ and R¹⁴, together with the carbon to which they are attached, form a spirocycloalkyl group of 3 to 6 carbons, or a
 10 spiroheterocycloalkyl group of 3 to 6 members, comprised of 2 to 5 carbon atoms and 1 or 2 heteroatoms selected from the group consisting of O, S and N; and

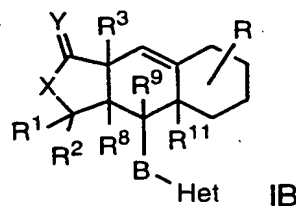
R²² is H, C₁-C₆ alkyl, phenyl, benzyl, -COR¹⁶ or -CONR¹⁸R¹⁹.

15 One group of preferred compounds is that of the formula IA



wherein X, Y, R, R¹, R², R³, R⁸, R⁹, R¹⁰, R¹¹, B and Het are as defined above.

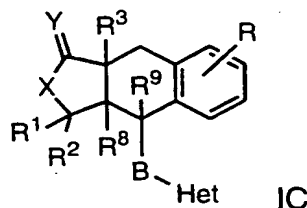
A second group of preferred compounds is that of the formula IB



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wherein X, Y, R, R¹, R², R³, R⁸, R⁹, R¹¹, B and Het are as defined above.

A third group of preferred compounds is that of the formula IC



wherein X, Y, R, R¹, R², R³, R⁸, R⁹, B and Het are as defined above.

R², R⁸, R¹⁰ and R¹¹ are each preferably hydrogen. R³ preferably is hydrogen or lower alkyl. The variable n is preferably zero. R⁹ is preferably H, OH or alkoxy. R¹ is preferably C₁-C₆ alkyl, more preferably methyl. The double dotted line preferably represents a single bond; X is preferably -O- and Y is preferably =O or (H, -OH). Z is preferably -CH₂-, -C(O)- or -C(=NOR¹⁷)-. The ring Q is preferably R-substituted cyclohexyl or R-substituted phenyl. R is preferably hydrogen, fluorine, hydroxy, alkoxy or alkyl. B is preferably trans -CH=CH-. Het is preferably pyridyl, substituted pyridyl, quinolyl or substituted quinolyl.

5 Preferred substituents (W) on Het are aryl, substituted aryl, heteroaryl or alkyl. More preferred are compounds wherein Het is 2-pyridyl substituted in the 5-position by aryl, substituted aryl, heteroaryl or alkyl, or 2-pyridyl substituted in the 6-position by alkyl.

Thrombin receptor antagonist compounds of the present invention have anti-thrombotic, anti-platelet aggregation, antiatherosclerotic, antirestenotic and anti-coagulant activity. Thrombosis-related diseases treated by the compounds of this invention are thrombosis, atherosclerosis, restenosis, hypertension, angina pectoris, arrhythmia, heart failure, myocardial infarction, glomerulonephritis, thrombotic and thromboembolytic stroke, peripheral vascular diseases, other cardiovascular diseases, cerebral ischemia, inflammatory disorders and cancer, as well as other disorders in which thrombin and its receptor play a pathological role.

15 20

Therefore, this invention also relates to the use of a compound of formula I as an antithrombotic, anti-platelet aggregation, anticoagulant or anticancer agent in a mammal in need of such treatment.

25

In another aspect, the invention relates to a pharmaceutical composition comprising a compound of formula I in a pharmaceutically acceptable carrier.

30

DETAILED DESCRIPTION:

Unless otherwise defined, the term "alkyl" or "lower alkyl" means straight or branched alkyl chains of 1 to 6 carbon atoms and "alkoxy" similarly refers to alkoxy groups having 1 to 6 carbon atoms.

35 Fluoroalkyl, difluoroalkyl and trifluoroalkyl mean alkyl chains wherein the terminal carbon is substituted by 1, 2 or 3 fluoroatoms, e.g.,

-CF₃, -CH₂CF₃, -CH₂CHF₂ or -CH₂CH₂F. Haloalkyl means an alkyl chain substituted by 1 to 3 halo atoms.

"Alkenyl" means straight or branched carbon chains of 1 to 6 carbon atoms having one or more double bonds in the chain, conjugated or unconjugated. Similarly, "alkynyl" means straight or branched carbon chains of 1 to 6 carbon atoms having one or more triple bonds in the chain. Where an alkyl, alkenyl or alkynyl chain joins two other variables and is therefore bivalent, the terms alkylene, alkenylene and alkynylene are used.

10 "Cycloalkyl" means a saturated carbon ring of 3 to 6 carbon atoms, while "cycloalkylene" refers to a corresponding bivalent ring, wherein the points of attachment to other groups include all positional and stereoisomers.

"Heterocycloalkyl" as a substituent on Het means saturated rings of 4 to 7 atoms comprised of 3 to 4 carbon atoms and 1 to 3 heteroatoms selected from the group consisting of -O-, -S- and -NR⁷- joined to the rest of the molecule through a carbon atom. Examples of heterocycloalkyl groups are 2-azetidiny, 2-pyrrolidinyl, tetrahydrothiophen-2-yl, tetrahydro-2-furanyl, 4-piperidinyl, 2-piperazinyl, tetrahydro-4-pyranyl, 2-morpholinyl and 2-thiomorpholinyl.

"Halogen" refers to fluorine, chlorine, bromine or iodine radicals.

When R⁴ and R⁵ join to form a ring with the nitrogen to which they are attached, the rings formed are 1-pyrrolidinyl, 1-piperidinyl and 1-piperazinyl, wherein the piperazinyl ring may also be substituted at the 4-position nitrogen by a group R⁷.

"Dihydroxy(C₁-C₆)alkyl" refers to an alkyl chain substituted by two hydroxy groups on two different carbon atoms.

"Aryl" means phenyl, naphthyl, indenyl, tetrahydronaphthyl or indanyl.

30 "Heteroaryl" means a single ring, bicyclic or benzofused heteroaromatic group of 5 to 10 atoms comprised of 2 to 9 carbon atoms and 1 to 4 heteroatoms independently selected from the group consisting of N, O and S, provided that the rings do not include adjacent oxygen and/or sulfur atoms. N-oxides of the ring nitrogens are also included, as well as compounds wherein a ring nitrogen is substituted by a C₁-C₄ alkyl group to form a quaternary amine. Examples of single-

ring heteroaryl groups are pyridyl, oxazolyl, isoxazolyl, oxadiazolyl, furanyl, pyrrolyl, thienyl, imidazolyl, pyrazolyl, tetrazolyl, thiazolyl, isothiazolyl, thiadiazolyl, pyrazinyl, pyrimidyl, pyridazinyl and triazolyl. Examples of bicyclic heteroaryl groups are naphthyridyl (e.g., 1, 5 or 1,7), imidazopyridyl, pyrido[2,3]imidazolyl, pyridopyrimidinyl and 7-azaindolyl. Examples of benzofused heteroaryl groups are indolyl, quinolyl, isoquinolyl, phthalazinyl, benzothienyl (i.e., thionaphthenyl), benzimidazolyl, benzofuranyl, benzoxazolyl and benzofurazanyl. All positional isomers are contemplated, e.g., 1-pyridyl, 2-pyridyl, 3-pyridyl and 4-pyridyl. W-substituted heteroaryl refers to such groups wherein substitutable ring carbon atoms have a substituent as defined above, or where adjacent carbon atoms form a ring with an alkylene group or a methylenedioxy group.

The term "Het" is exemplified by the single ring, bicyclic and benzofused heteroaryl groups as defined immediately above, as well as tricyclic groups such as benzoquinolinyl (e.g., 1,4 or 7,8) or phenanthrolinyl (e.g., 1,7; 1,10; or 4,7). Het groups are joined to group B by a carbon ring member, e.g., Het is 2-pyridyl, 3-pyridyl or 2-quinolyl.

Examples of heteroaryl groups wherein adjacent carbon atoms form a ring with an alkylene group are 2,3-cyclopentenopyridine, 2,3-cyclohexenopyridine and 2,3-cycloheptenopyridine.

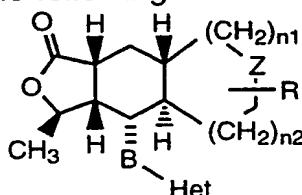
The term "optional double bond" refers to the bond shown by the single dotted line in the middle ring of the structure shown for formula I. The term "optional single bond" refers to the bond shown by the double dotted line between X and the carbon to which Y and R¹⁵ are attached in the structure of formula I. Compounds wherein the bond is absent are shown, for example, in Example 6.

The above statements, wherein, for example, R⁴ and R⁵ are said to be independently selected from a group of substituents, means that R⁴ and R⁵ are independently selected, but also that where an R⁴ or R⁵ variable occurs more than once in a molecule, those occurrences are independently selected. Those skilled in the art will recognize that the size and nature of the substituent(s) will affect the number of substituents which can be present.

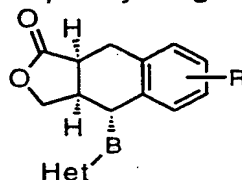
Compounds of the invention have at least one asymmetrical carbon atom and therefore all isomers, including diastereomers and

- rotational isomers are contemplated as being part of this invention. The invention includes (+)- and (-)-isomers in both pure form and in admixture, including racemic mixtures. Isomers can be prepared using conventional techniques, either by reacting optically pure or optically enriched starting materials or by separating isomers of a compound of formula I.

Typical preferred compounds of the present invention wherein Q is a saturated ring have the following stereochemistry:



- with compounds having that absolute stereochemistry being more preferred. Typical preferred compounds of the present invention wherein Q is an aromatic ring have the following stereochemistry, wherein Q is exemplified as a phenyl ring:



- with compounds having that absolute stereochemistry being more preferred.

Those skilled in the art will appreciate that for some compounds of formula I, one isomer will show greater pharmacological activity than other isomers.

- Compounds of the invention with a basic group can form pharmaceutically acceptable salts with organic and inorganic acids. Examples of suitable acids for salt formation are hydrochloric, sulfuric, phosphoric, acetic, citric, oxalic, malonic, salicylic, malic, fumaric, succinic, ascorbic, maleic, methanesulfonic and other mineral and carboxylic acids well known to those in the art. The salt is prepared by contacting the free base form with a sufficient amount of the desired acid to produce a salt. The free base form may be regenerated by treating the salt with a suitable dilute aqueous base solution such as dilute aqueous sodium bicarbonate. The free base form differs from its

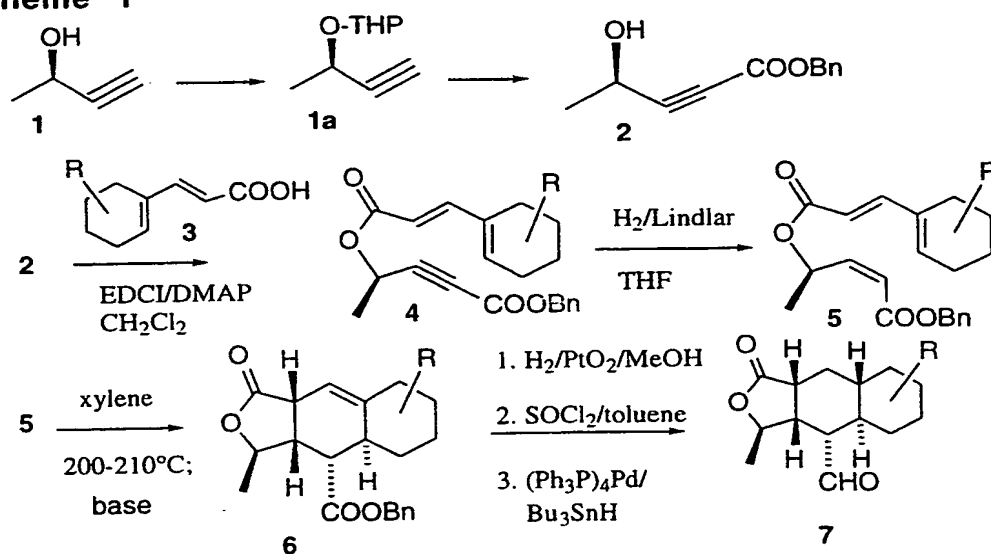
respective salt form somewhat in certain physical properties, such as solubility in polar solvents, but the salt is otherwise equivalent to its respective free base forms for purposes of the invention.

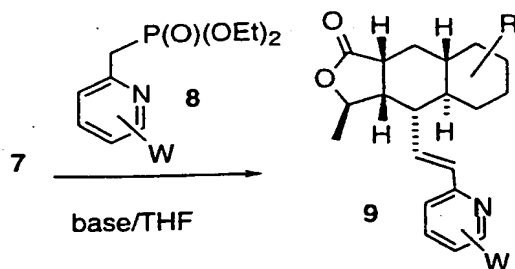
Certain compounds of the invention are acidic (e.g., those compounds which possess a carboxyl group). These compounds form pharmaceutically acceptable salts with inorganic and organic bases. Examples of such salts are the sodium, potassium, calcium, aluminum, lithium, gold and silver salts. Also included are salts formed with pharmaceutically acceptable amines such as ammonia, alkyl amines, hydroxyalkylamines, N-methylglucamine and the like.

Compounds of the present invention are generally prepared by processes known in the art, for example by the processes described below.

In Scheme 1, a process is shown for preparing compounds of formula I wherein n is 0, the optional double bond is not present, Q forms a cyclohexyl ring, the single bond is present between X and the carbon to which Y is attached, X is -O-, Y is =O, B is -CH=CH-, Het is W-substituted pyridyl, R² is methyl, and R¹, R³, R⁸, R⁹, R¹⁰ and R¹¹ are each hydrogen. However, a similar process may be used to prepare compounds comprising other optionally substituted Het groups. Those skilled in the art will also recognize that the process is equally applicable to preparing optically active or racemic compounds.

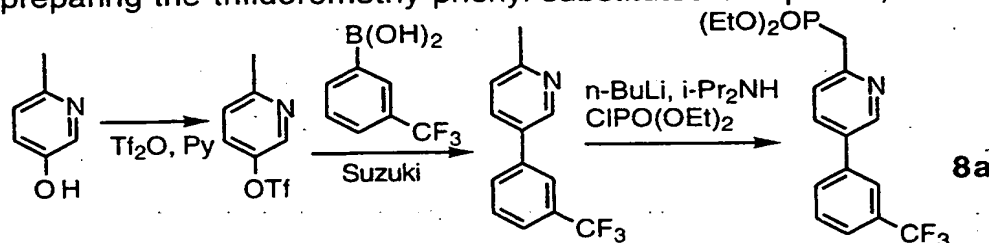
Scheme 1





- Commercially available (*R*)-3-butyn-2-ol is *O*-protected as the tetrahydropyranyl ether by treatment with dihydropyran in the presence of catalytic amounts of para-toluenesulfonic acid to give intermediate
- 5 **1a**. Treatment of a solution of **1a** in THF with *n*-BuLi at -78°C followed by quenching with benzylchloroformate and subsequent deprotection gives intermediate **2** which is esterified with dienoic acid **3** using standard conditions to yield the ester **4**. Selective reduction of the triple bond of **4** using Lindlar catalyst under hydrogen gives the intermediate
- 10 **5**, which upon thermal cyclization between 200-210°C followed by base treatment gives the tricyclic intermediate **6**. The ester **6** is subjected to hydrogenation in the presence of platinum oxide to generate the intermediate saturated carboxylic acid, treatment of which with SOCl₂ gives the corresponding acid chloride which is converted to the tricyclic
- 15 aldehyde **7** by reduction using tributyltin hydride in the presence of Pd(0). Condensation of the anion generated from phosphonate **8** with aldehyde **7** in THF gives the alkene **9** (final product).

- Intermediates of formula **8** wherein W is aryl or R²¹-aryl can be prepared by a process similar to that described immediately below for preparing the trifluoromethyl-phenyl-substituted compound, **8a**.
- 20



- Commercially available hydroxypyridine derivative is converted to the corresponding triflate using triflic anhydride, which is then coupled with commercially available boronic acid in the presence of Pd(0) under
- 25 Suzuki conditions. The resulting product is converted to the

phosphonate by treatment with tert-butyllithium followed by quenching with diethylchlorophosphonate.

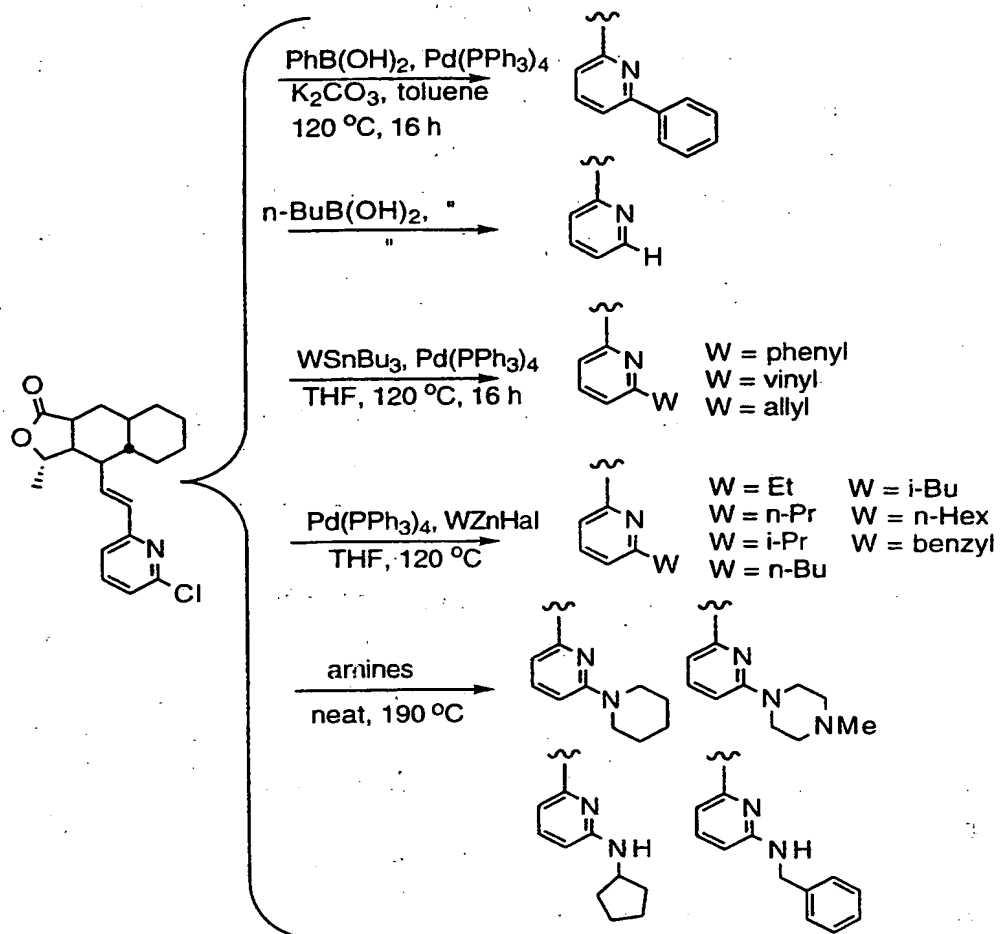
Alternatively, compounds of formula 9 wherein W is optionally substituted aryl can be prepared from compounds of formula 9 wherein W is -OH using a triflate intermediate. For example, 3-hydroxy-6-methylpyridine is treated with triisopropylsilyl chloride, and the resultant hydroxy-protected compound is converted to the phosphonate as described above for preparing intermediate 8. The triisopropylsilyl-protected intermediate is then reacted with tricyclic intermediate 7 and the protecting group is removed under standard conditions. The resultant compound of formula 9 wherein W is OH is then treated with triflic anhydride at room temperature in a solvent such as CH₂Cl₂; the triflate is then reacted with an optionally substituted arylboronic acid, e.g., optionally substituted phenylboronic acid, in a solvent such as toluene, in the presence of Pd(PPh₃)₄ and a base such as K₂CO₃ at elevated temperatures and under an inert atmosphere.

Compounds of formula 9 wherein W is a substituted hydroxy group (e.g., benzyloxy) can be prepared from compounds of formula 9 wherein W is hydroxy by refluxing in a suitable solvent such as acetone with a halogen-substituted compound such as optionally substituted benzyl bromide in the presence of a base such as K₂CO₃.

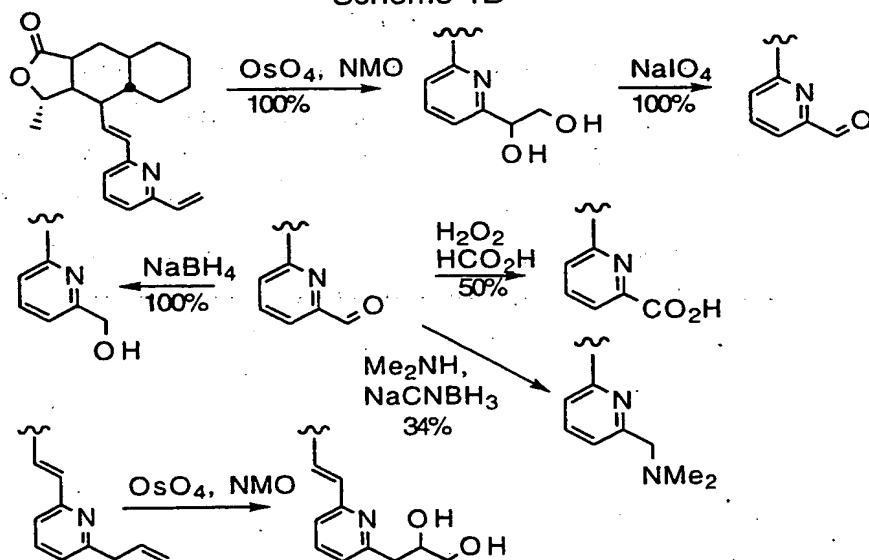
Compounds of formula I wherein Het is substituted by W through a carbon atom (e.g., wherein W is alkyl, alkenyl or arylalkyl) or a nitrogen atom (i.e., -NR⁴R⁵) can be prepared using a compound of formula I wherein W is chloroalkyl as an intermediate. Compounds of formula I wherein W is a polar group such as hydroxy alkyl, dihydroxyalkyl, -COOH, dimethylamino and -COH can be prepared as shown in Scheme 1B, wherein the starting material is a compound of formula I wherein W is alkenyl. The following Schemes 1A and 1B show well-known reaction conditions for preparing various W-substituted compounds wherein Q is cyclohexyl, X is -O-, Y is =O, R¹⁵ is absent, R¹ is methyl, R², R³, R⁹, R¹⁰ and R¹¹ are each H, B is -CH=CH-, and Het is 2-pyridyl.

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Scheme 1A



Scheme 1B



Those skilled in the art will appreciate that similar reactions to those described in the above tables may be carried out on other compounds of formula I as long as substituents present would not be susceptible to the reaction conditions described.

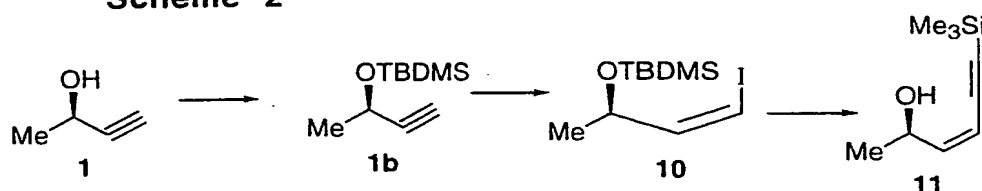
- 5 Compounds of formula I wherein Q is non-aromatic, the optional single bond (represented by the double dotted line) is absent, X is OH, Y is OH, R¹⁵ is H and the remaining variables are as defined above can be prepared by treating corresponding compounds wherein the optional single bond is present, X is -O-, Y is =O and R¹⁵ is absent, with a
10 reducing agent such as LAH.

- Compounds of formula I wherein the optional single bond is present, X is -O-, Y is (H, OH), R¹⁵ is absent and the remaining variables are as defined above can be prepared by treating corresponding compounds wherein the optional single bond is present, X is -O-, Y is =O
15 and R¹⁵ is absent, with a reagent such as DIBAL. The resultant compounds wherein Y is (H, OH) can be converted to the corresponding compounds wherein Y is (H, alkoxy) by reacting the hydroxy compound with an appropriate alkanol in the presence of a reagent such as BF₃•OEt₂. A compound wherein Y is (H, OH) can also be converted to
20 the corresponding compound wherein Y is (H, H) by treating the hydroxy compound with BF₃•OEt₂ and Et₃SiH in an inert solvent such as CH₂Cl₂ at low temperatures.

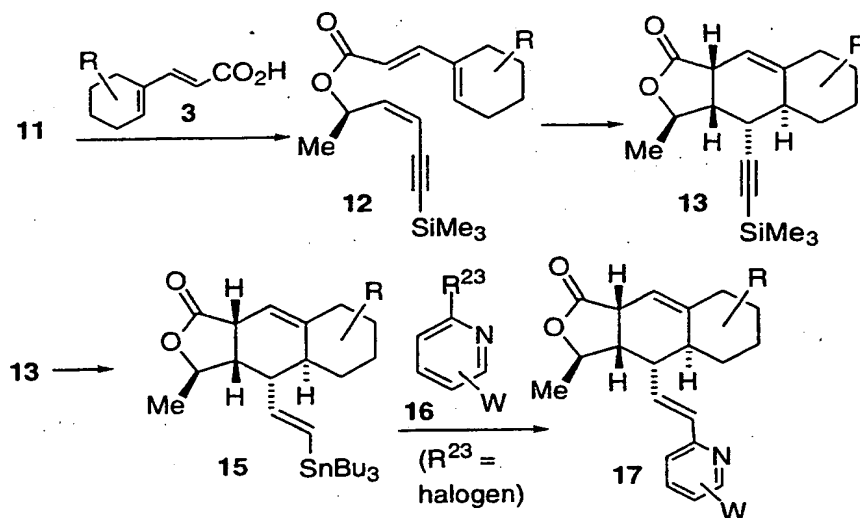
- Compounds of formula I wherein R⁹ is hydrogen can be converted to the corresponding compound wherein R⁹ is hydroxy by
25 heating with an oxidizing agent such as SeO₂.

- In Scheme 2, a process is shown for preparing compounds of formula I wherein n is 0, the optional double bond is present, Q forms a cyclohexyl ring, X is -O-, Y is =O, R², R³, R⁸, R⁹ and R¹¹ are each hydrogen, R¹ is methyl, R¹⁰ is absent, B is -CH=CH-, and Het is W-substituted pyridyl. However, a similar process may be used to prepare
30 compounds comprising other optionally substituted Het groups.

Scheme 2

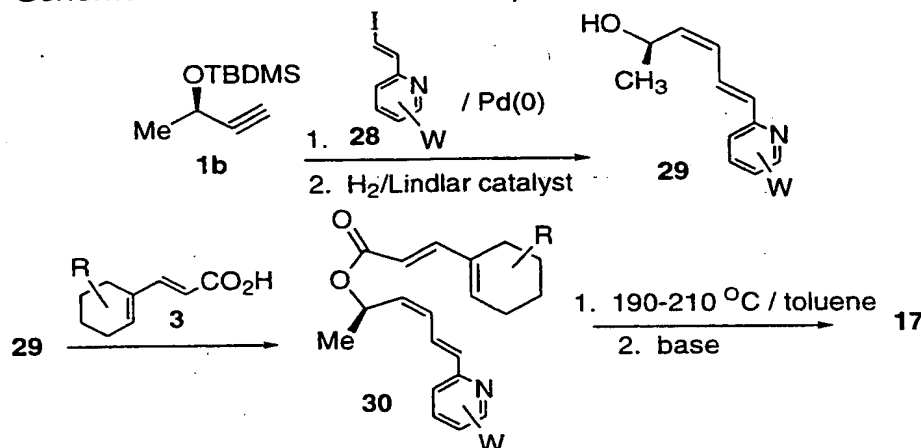


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The alcohol **1** is O-protected as the TBDMS ether by treatment with TBDMS chloride. The anion generated from **1b** is quenched with a solution of iodine to give the corresponding acetylenic iodide, which upon reduction using di(cyclohexyl)borane gives the cis-vinyl iodide **10**. Coupling of vinyl iodide **10** with trimethylsilylacetylene in the presence of Cu(I) and Pd(0) gives the intermediate **11** after removal of the protecting group using trifluoroacetic acid in CH₃OH. Esterification of **11** with acid **3** gives the intermediate ester **12**, which upon thermal cyclization at 185-195°C gives the tricyclic precursor **13** after a brief treatment with DBU. Desilylation of acetylenic derivative **13** followed by hydrostannylation using tributyltinhydride in the presence of AIBN gives the vinylstannane derivative **15**, which is coupled with halopyridine derivative **16** to give the final product **17**.

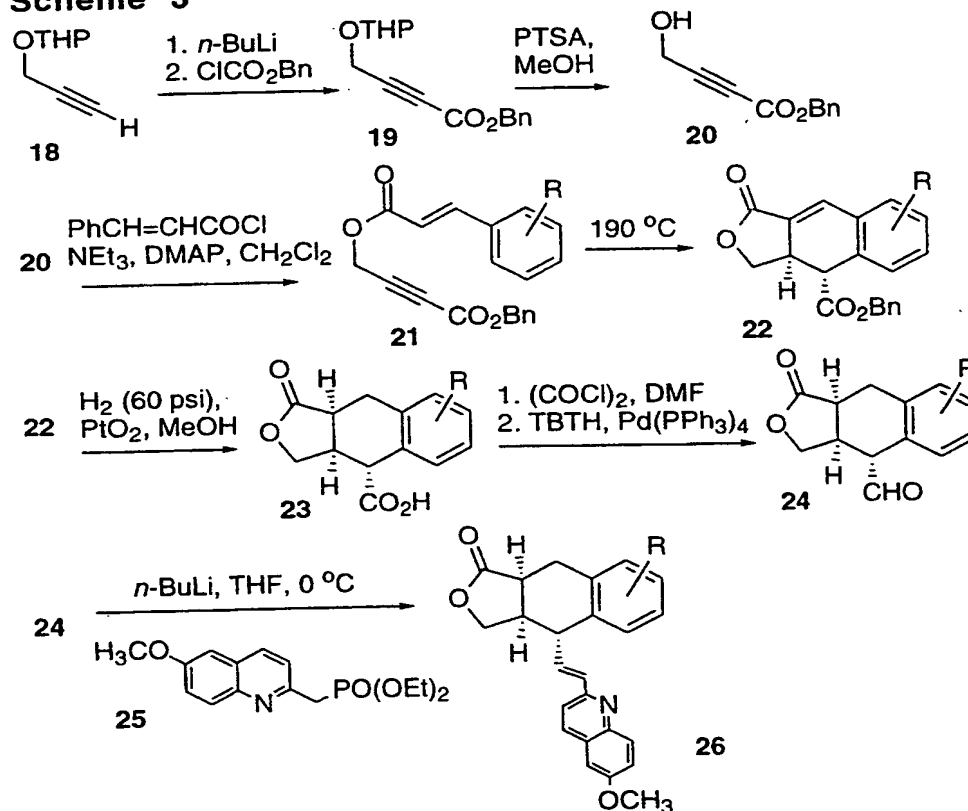
Scheme 2A offers an alternative procedure:



Palladium-mediated coupling of acetylene **1b** with trans-iodovinylpyridine **28** followed by selective reduction of the triple bond gives the intermediate dienoic alcohol **29** which is esterified with the dienoic acid **3** to give **30**. Thermal cyclization of **30** at 190-210°C followed by base treatment gives **17**. Intermediate **28** is prepared from (2-chloro-6-methyl)pyridine by coupling with (trimethylsilyl)acetylene in the presence of palladium, followed by deprotection of the silyl group using fluoride anion and treatment of the isolated product with tributyltin hydride followed by iodine.

In Scheme 3, a process is shown for preparing compounds of formula I wherein n is 0, the optional double bond is not present, Q forms a phenyl ring, X is -O-, Y is =O, R¹, R², R³, R⁸ and R⁹ are each hydrogen, R¹⁰ and R¹¹ are absent, B is -CH=CH-, and Het is methoxy-substituted quinolyl. However, a similar process may be used to prepare compounds comprising other optionally substituted Het groups.

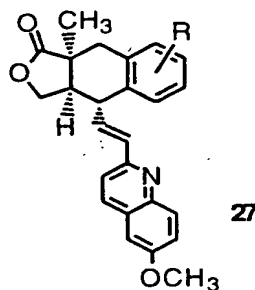
Scheme 3



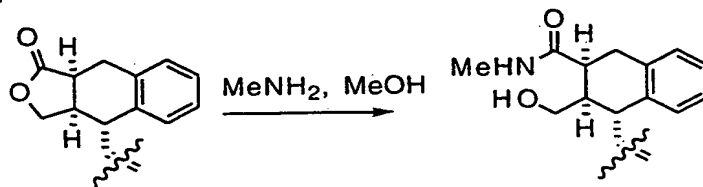
Commercially available acetylenic derivative **18** is converted to the corresponding ester **19** by treatment with *n*-BuLi in THF followed by

quenching with benzylchloroformate. Removal of the THP group followed by esterification using *trans*-cinnamyl chloride under standard conditions gives the ester **21** which, upon thermal cyclization at 190°C, gives the tricyclic ester **22**. Hydrogenation of **22** over platinum oxide gives the carboxylic acid **23** which is converted to the corresponding acid chloride under standard conditions. Reduction of the acid chloride generated from **23** using Pd(0) and tributyltin hydride gives the aldehyde **24** which is condensed with the anion generated from the phosphonate **25** to give the final product, **26**.

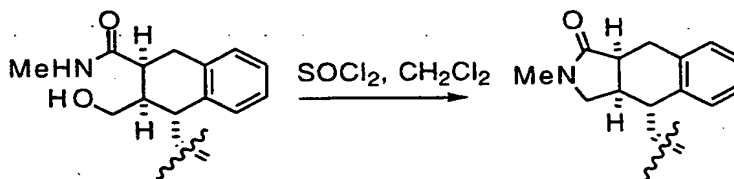
Compounds wherein R³ is alkyl can be prepared from the corresponding compounds wherein R³ is hydrogen. For example, treatment of compound **26** with LDA followed by CH₃I results in the preparation of the corresponding compound **27** wherein R³ is methyl:



Amide, lactam and imide derivatives of compounds of formula I wherein Q is an aromatic ring can be prepared from compounds wherein X is -O- and Y is =O using the following procedures showing partial structural formulas:

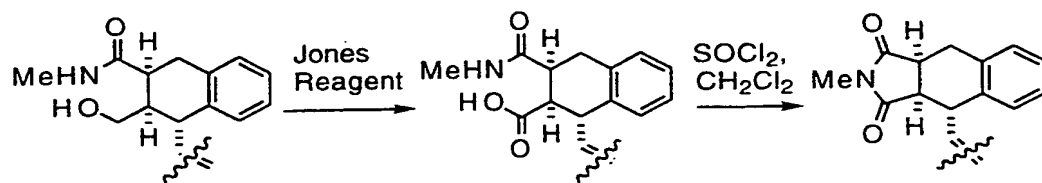


The lactone is treated with an amine to form the ring-opened amide.



The amide is treated with a reagent such as SOCl₂ to cyclize the amide to the lactam.

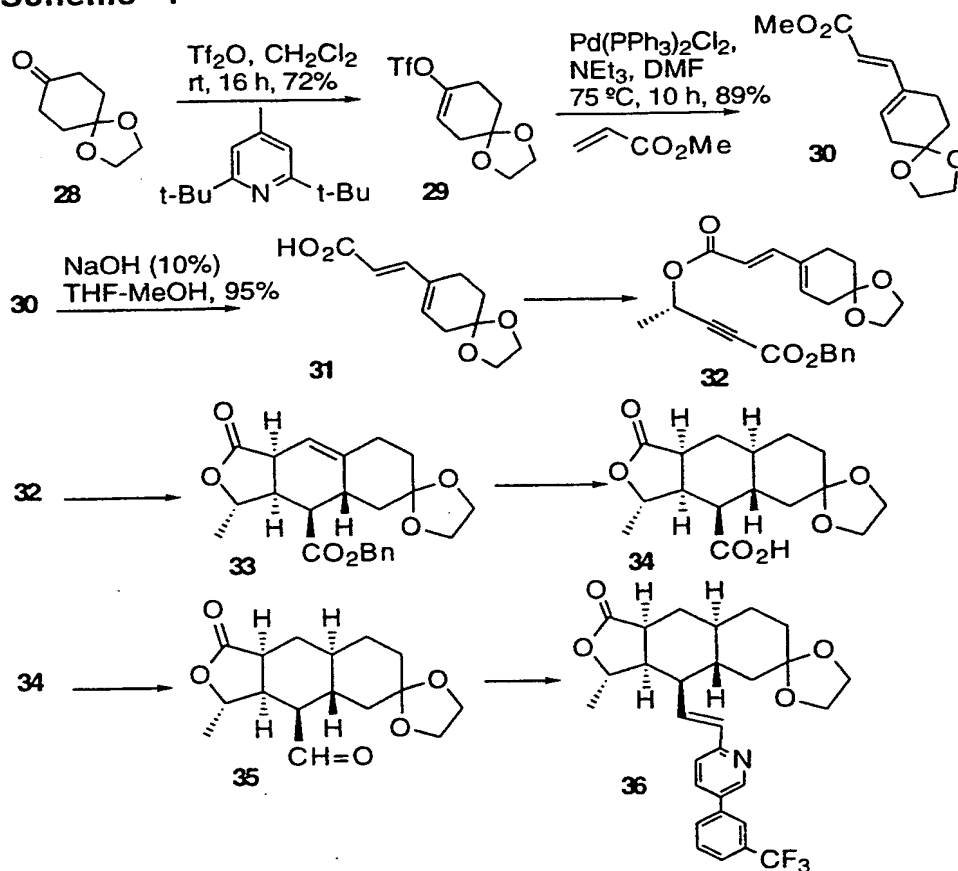
-18-



The hydroxy-substituted portion of the amide can be converted to a carboxylic acid by treatment with a reagent such as Jones Reagent, and the resultant product is cyclized to form the imide in the same manner as for the lactam.

In Scheme 4, a process is shown for preparing compounds of formula I wherein n is 0, the optional double bond is not present, Q is a cyclohexyl ring substituted in the 6-position by ethylenedioxy (i.e., Z is -C(R¹³R¹⁴)), X is -O-, Y is =O, R¹, R³, R⁸, R⁹, R¹⁰ and R¹¹ are each hydrogen, R² is methyl, R¹⁵ is absent, B is -CH=CH-, Het is pyridyl and W is CF₃-phenyl. However, a similar process may be used to prepare compounds comprising other optionally substituted Het groups.

Scheme 4

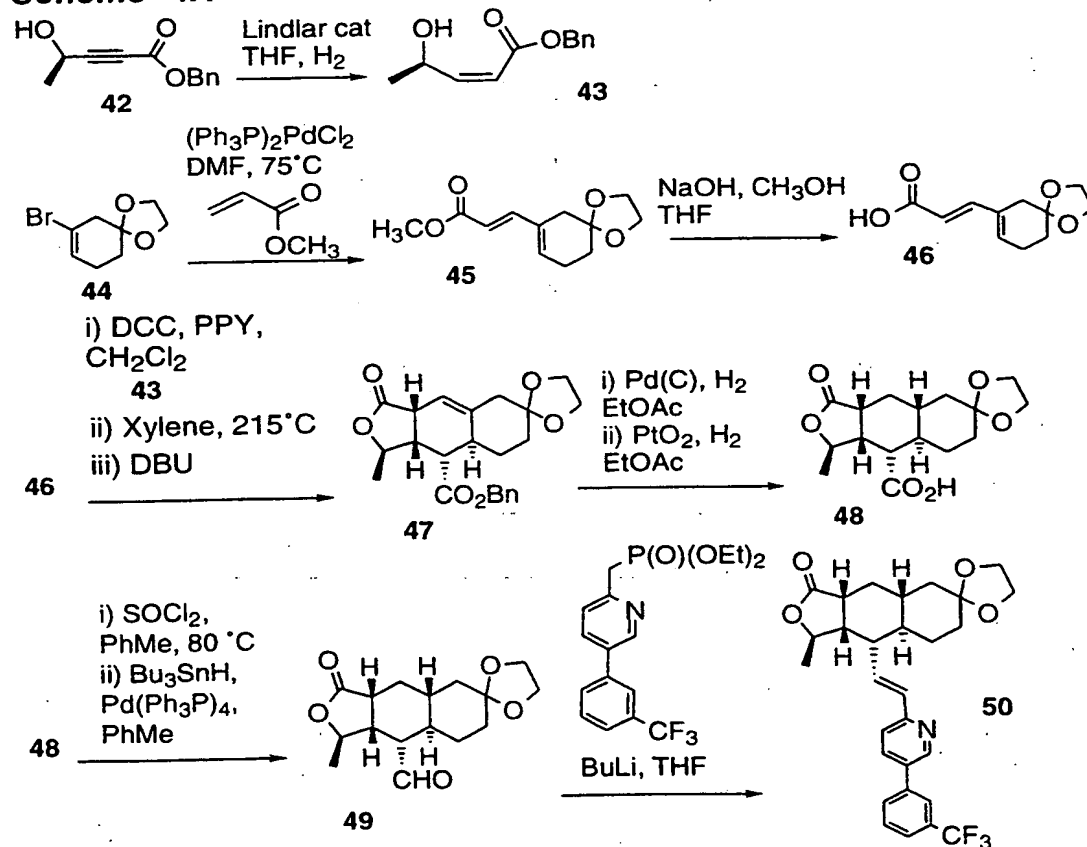


A solution of 1,4-cyclohexanedione mono-ethylene ketal, **28**, and 2,6-di-*t*-butyl-4-methylpyridine is treated with triflic anhydride to obtain the enol triflate, **29**. Compound **29** is converted to compound **30** by treatment with methyl acrylate in a solvent such as DMF, in the presence of a base such as Et₃N and a catalyst such as Pd(PPh₃)₂Cl₂, and **30** is converted to the corresponding acid, **31**, by standard methods, e.g. by treatment with a base such as NaOH. Acid **31** is then reacted with racemic **2** and cyclized as described in Scheme 1 to obtain the product **36** as a racemic mixture.

A ketal such as compound **36** can be converted to the corresponding ketone by treatment with an acid such as HCl. The ketone, in turn, can be reduced to the corresponding hydroxy compound by treatment with a reagent such as NaBH₄ or K-Selectride®.

In Scheme 4A, a process is shown for preparing compounds similar to those in Scheme 4, but wherein Z is ethylenedioxy and is in the 7-position of the cyclohexyl ring.

Scheme 4A



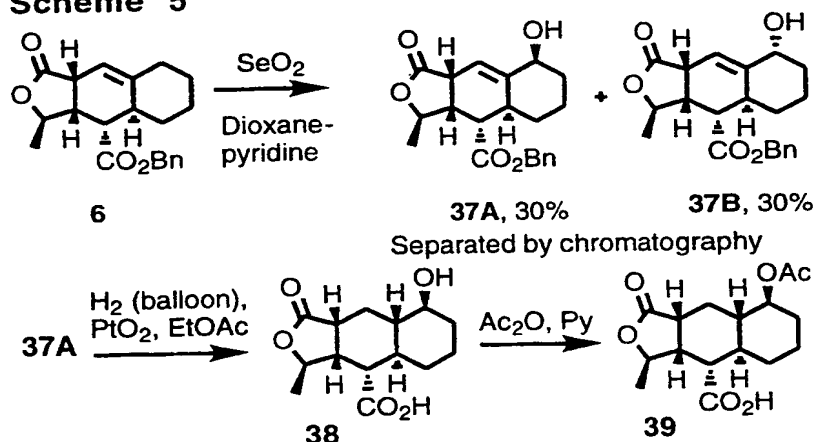
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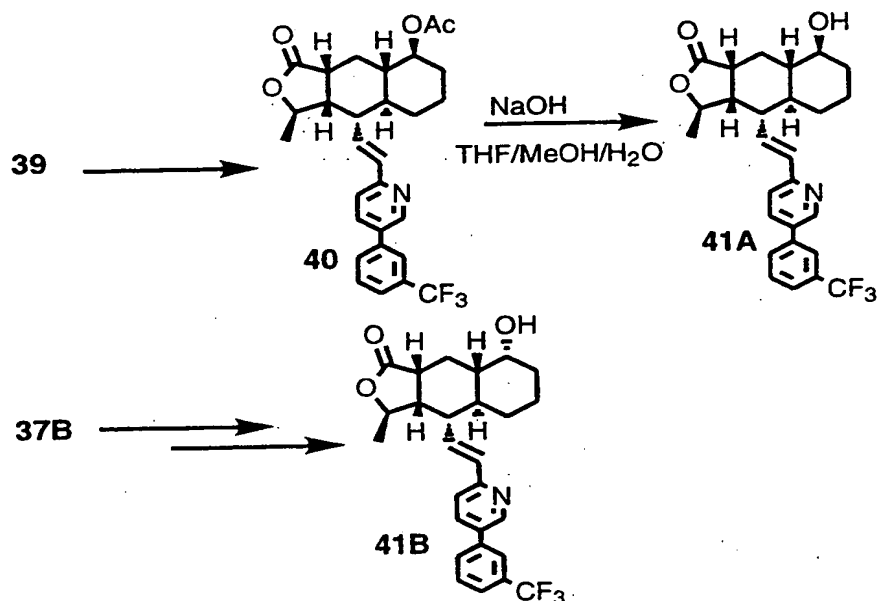
A solution of the ketal, **44**, is treated with methyl acrylate in a solvent such as DMF, in the presence of a base such as Et₃N and a catalyst such as Pd(PPh₃)₂Cl₂, and the resultant ester, **45**, is converted to the corresponding acid, **46**, by standard methods, e.g. by treatment
 5 with a base such as NaOH. Acid **46** is then reacted with compound **43** and cyclized to obtain **47**, which is then converted to the corresponding acid by standard methods. The intermediate of **49** is coupled to the -B-Het group using a procedure as described in Scheme 1 to obtain compound **50**.

10 A ketal such as compound **50** can be converted to the corresponding ketone by treatment with an acid such as HCl. The ketone, in turn, can be reduced to the corresponding hydroxy compound by treatment with a reagent such as NaBH₄ or K-Selectride[®]. The
 15 ketone can be converted to the corresponding 7-hydroxy-7 methyl compound by treatment with a reagent such as CH₃MgBr.

In Scheme 5, a process is shown for preparing compounds of formula I wherein n is 0, the optional double bond is absent, Q forms a cyclohexyl ring, X is -O-, Y is =O, R¹, R³, R⁸, R⁹, R¹⁰ and R¹¹ are each H, R² is -CH₃, R¹⁵ is absent, B is -CH=CH-, Het is pyridyl, W is CF₃-phenyl,
 20 and R is hydroxy. However, a similar process may be used to prepare compounds comprising other optionally substituted Het groups.

Scheme 5





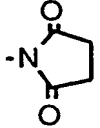
Intermediate 6 from Scheme 1 is oxidized to give intermediate alcohols **37A** and **37B**. Alcohol **37A** is hydrogenated to give **38** which is subsequently converted to acetate **39**. Acetate **39** is converted to intermediate **40** in the same fashion as in Scheme 1. Intermediate **40** is hydrolyzed to give **41A**. Using a similar procedure, but substituting **37B** in the second step, a compound of formula **41B** is obtained.

Compounds of formula I wherein B is $-\text{C}(\text{O})\text{NH}-$ can be prepared from the acid chloride described above (see Scheme 3) by condensation with the appropriate amine.

Starting materials for the above processes are either commercially available, known in the art, or prepared by procedures well known in the art.

Reactive groups not involved in the above processes can be protected during the reactions with conventional protecting groups which can be removed by standard procedures after the reaction. The following Table A shows some typical protecting groups:

Table A

Group to be Protected	Group to be Protected and Protecting Group
-COOH	-COOalkyl, -COObenzyl, -COOphenyl
>NH	$\begin{array}{l} \diagup \text{NCOalkyl} \diagdown, \diagup \text{NCObenzyl} \diagdown, \diagup \text{NCOphenyl} \diagdown \\ \diagdown \text{NCH}_2\text{OCH}_2\text{CH}_2\text{Si}(\text{CH}_3)_3 \diagup, \diagdown \text{NC}(\text{O})\text{OC}(\text{CH}_3)_3 \diagup \\ \diagdown \text{N-benzyl} \diagup, \diagdown \text{NSi}(\text{CH}_3)_3 \diagup, \begin{array}{c} \text{CH}_3 \\ \\ \diagdown \text{NSi-C}(\text{CH}_3)_3 \diagup \\ \\ \text{CH}_3 \end{array} \end{array}$
-NH ₂	
-OH	$\begin{array}{l} -\text{OCH}_3, -\text{OCH}_2\text{OCH}_3, -\text{OSi}(\text{CH}_3)_3, -\begin{array}{c} \text{CH}_3 \\ \\ \text{OSi-C}(\text{CH}_3)_3 \\ \\ \text{CH}_3 \end{array} \\ \text{or } -\text{OCH}_2\text{phenyl} \end{array}$

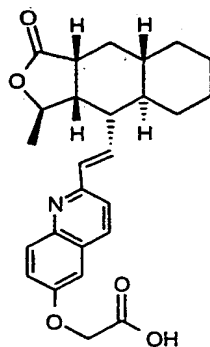
The present invention also relates to a pharmaceutical composition comprising a compound of formula I of this invention and a pharmaceutically acceptable carrier. The compounds of formula I can be administered in any conventional oral dosage form such as capsules, tablets, powders, cachets, suspensions or solutions. The formulations and pharmaceutical compositions can be prepared using conventional pharmaceutically acceptable excipients and additives and conventional techniques. Such pharmaceutically acceptable excipients and additives include non-toxic compatible fillers, binders, disintegrants, buffers, preservatives, anti-oxidants, lubricants, flavorings, thickeners, coloring agents, emulsifiers and the like.

The daily dose of a compound of formula I for treatment of a disease or condition cited above is about 0.001 to about 100 mg/kg of body weight per day, preferably about 0.001 to about 10 mg/kg. For an average body weight of 70kg, the dosage level is therefore from about 0.1 to about 700 mg of drug per day, given in a single dose or 2-4 divided doses. The exact dose, however, is determined by the attending clinician and is dependent on the potency of the compound administered, the age, weight, condition and response of the patient.

Following are examples of preparing compounds of formula I.

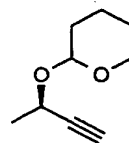
EXAMPLE 1

[[2-[(E)-2-(3R, 3aS, 4S, 8aS, 9aR-Dodecahydro-3-methyl-1-oxo-naphtho[2,3-c]furan-4-yl)ethenyl]-6-quinolinyloxy]acetic acid



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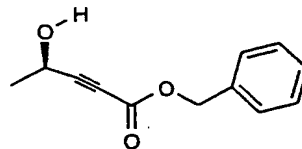
Step 1:



(R)-3-Butyn-2-ol (15 ml, 0.204 mol) and 3,4-dihydro-2H-pyran (26.1 ml, 1 eq) were stirred at 0°C. *p*-Toluenesulfonic acid(hydrate) (0.38 g, 5 mol %) was added and the mixture stirred for a further 2 h. Ethyl Acetate (EtOAc) (319 ml) and NaHCO₃ (1.6 g) were added and after 1 h the mixture was filtered and concentrated. Chromatography (SiO₂, 19:1 hexane/EtOAc) gave 31.49 g (100%) of the desired product as a mixture of diastereomers. ¹H NMR *major diastereomer* (CDCl₃) δ 1.54 (d, *J* = 7.5 Hz, 3H), 1.55-2.0 (m, 6H), 2.42 (s, 1H), 3.56 (m, 1H), 3.88 (m, 1H), 4.60 (br q, *J* = 7.5 Hz, 1H), 5.00 (t, *J* = 5.0 Hz, 1H).

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Step 2:



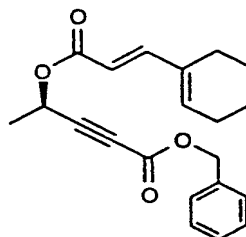
The product of Step 1 (31.49 g, 0.204 mol) was dissolved in THF (1 L) and cooled to -78°C with stirring. *n*-Butyllithium (97.8 ml of a 2.5 M solution, 1.2 eq) was added dropwise. After stirring for 20 min, benzyl chloroformate (35.1 ml, 1.2 eq) was added and the reaction stirred at -78°C for 2 h. The mixture was allowed to warm to room temperature (RT), NH₄Cl solution (saturated) was added and the mixture was

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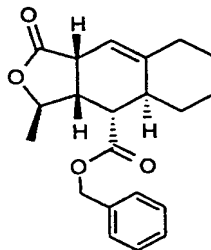
extracted with EtOAc. The organic extracts were dried (MgSO₄), concentrated and then dissolved in CH₃OH (2 L), DOWEX 50WX8-100 ion-exchange resin (60 g, prewashed with CH₃OH) was added and the mixture stirred at RT overnight. The mixture was filtered, concentrated and chromatographed (SiO₂, 9:1-4:1 hexane/EtOAc) to give 29.9 g (71 %) of the desired product. ¹H NMR (CDCl₃) δ 1.55 (d, *J* = 7.5 Hz, 3H), 4.70 (q, *J* = 7.5 Hz, 1H), 5.27 (s, 2H), 7.44 (br s, 5H).

Step 3:



trans-3-(1-Cyclohexenyl)acrylic acid (4.13 g, 0.0273 mol) and 4-pyrrolidinopyridine (0.4 g, 10 mol %) in CH₂Cl₂ (100 ml) were stirred at 0°C. 1,3 Dicyclohexylcarbodiimide (5.63 g, 1 eq) was added and the mixture stirred for 10 min. A solution of the product of Step 2 (5.58 g, 0.0273 mol) in CH₂Cl₂ (40 ml) was added dropwise. The resulting mixture was stirred for 2 hs, filtered, concentrated and chromatographed (SiO₂, 97:3 hexane/EtOAc) to give 5.82 g (63 %) of the desired product. ¹H NMR (CDCl₃) δ 1.61 (d, *J* = 7.0 Hz, 3H), 1.66 (m, 2H), 1.74 (m, 2H), 2.18 (m, 2H), 2.27 (m, 2H), 5.25 (s, 2H), 5.67 (m, 1H), 5.80 (d, *J* = 15.0 Hz, 1H), 6.26 (br s, 1H), 7.37 (d, *J* = 15.0 Hz, 1H), 7.42 (s, 5H).

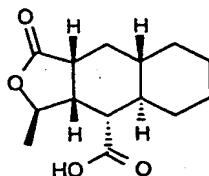
Step 4:



The product of Step 3 (5.82 g, 0.017 mol) and triethylamine (Et₃N) (0.112 mL) were dissolved in THF (32 mL). Lindlar catalyst (0.58 g) was added and the mixture stirred under 1 atm. of hydrogen for 16h. The mixture was filtered, concentrated, dissolved in o-xylene, and degassed under a stream of N₂. The degassed mixture was sealed in a pressure tube and heated at 210°C for 6 h. After cooling to RT, the xylene was

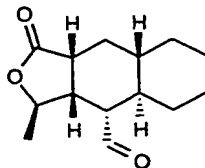
removed under reduced pressure and the resulting mixture chromatographed (SiO₂, 19:1-9:1 hexane/EtOAc) to give 3.81 g (66 %) of the desired product. ¹H NMR (CDCl₃) δ 0.94 (m, 1H), 1.20 (d, J = 7.0 Hz, 3H), 1.31 (m, 1H), 1.50 (m, 1H), 1.82 (m, 2H), 2.00 (m, 1H), 2.20 (m, 1H), 2.39 (br d, J = 15.0 Hz, 1H), 2.51 (m, 1H), 2.62 (m, 1H), 2.73 (m, 1H), 3.35 (m, 1H), 4.53 (m, 1H), 5.22 (AB quartet, J = 12.5 Hz, 2H), 5.34 (br s, 1H), 7.42 (br s, 5H).

Step 5:



The product of Step 4 (3.81 g, 0.011 mol) was dissolved in CH₃OH (100 ml). Platinum (IV) oxide (0.38 g) was added and the mixture shaken for 16 h under an atmosphere of hydrogen (60 psi). The mixture was filtered, concentrated and recrystallized (CH₂Cl₂/ hexanes) to give 2.12 g (75 %) of the desired product. ¹H NMR (CDCl₃) δ 0.90-1.0 (m, 1H), 1.05-1.20 (m, 2H), 2.21-1.55 (m, 7H), 1.75-1.92 (m, 4H), 1.92-2.00 (m, 1H), 2.52-2.64 (m, 2H), 2.74 (m, 1H), 4.76 (m, 1H).

Step 6:

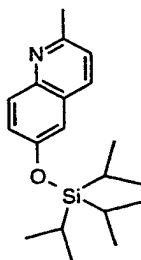


The product of Step 5 (2.3 g, 9.66 mmol) was suspended in toluene (20 ml), SOCl₂ (4 ml) was added and the mixture heated at 80°C for 16 h. After cooling to RT, the volatiles were removed under reduced pressure and the resulting gum dissolved in fresh toluene (23 ml). Tetrakis(triphenylphosphine)palladium(0) (800 mg, 8 mol %) was added and the mixture was cooled to 0°C. Tributyltin hydride (Bu₃SnH) (3.24 ml, 1.2 eq) was added dropwise and the mixture stirred for 30 min, after which time TLC showed approximately 66 % conversion. Bu₃SnH (1.35 mL, 0.5 eq) was added and the mixture stirred for a further hour. The mixture was then chromatographed (SiO₂, 4:1 hexanes/EtOAc) to give 1.9 g (88 %) of the desired product. ¹H NMR (CDCl₃) δ 0.88-1.05 (m, 1H), 1.10-1.20 (m, 2H), 1.22-1.50 (m, 5H), 1.55-1.70 (m, 2H), 1.75-

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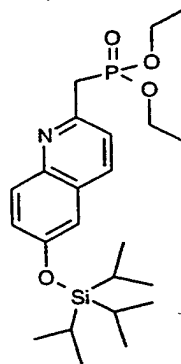
1.90 (m, 4H), 1.98 (dd, $J = 12.5, 7.0$ Hz, 1H) 2.53 (m, 1H), 2.63 (m, 1H), 2.73 (m, 1H), 4.73 (m, 1H), 9.80 (d, $J = 5.0$ Hz, 1H).

Step 7:



- 6-Hydroxyquinaldine (1.97 g, 0.0123 mol) and imidazole (0.85 g, 0.0124 mol) were dissolved in DMF (20 ml) and cooled to 0°C with stirring. Triisopropylsilyl chloride (2.7 ml, 1.05 eq) was added and the mixture stirred for 30 min. NH_4Cl solution (saturated) was added and the mixture extracted with EtOAc. The organic extracts were dried (MgSO_4), concentrated and chromatographed (SiO_2 , 4:1-1:1 hexane/EtOAc) to give 3.39 g (88 %) of the desired compound. ^1H NMR (CDCl_3) δ 1.12(d, $J = 8.5$ Hz, 18H), 1.30(heptet, $J = 8.5$ Hz, 2H), 2.05 (s, 3H), 7.13 (br s, 1H), 7.22 (d, $J = 9.0$ Hz, 1H), 7.31 (d, $J = 11.0$ Hz), 7.89 (m, 2H).

Step 8:

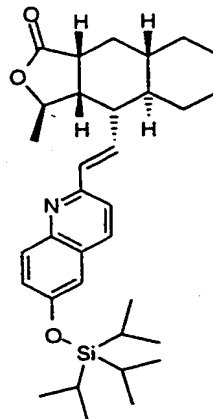


- The product of Step 6 (3.39 g, 0.0108 mol) and diisopropylamine (1.66 ml, 1 eq) were dissolved in THF (54 mL) and cooled to -78°C with stirring. *n*-Butyllithium (9 ml of a 2.5 M solution in hexanes, 2.1 eq) was added dropwise and after 20 min diethylchlorophosphate (1.7 ml, 1.1 eq) was added. After a further 20 min, the mixture was allowed to warm to RT. NH_4Cl solution (saturated) was added and the mixture extracted with EtOAc. The organic extracts were dried (MgSO_4), concentrated and chromatographed (SiO_2 , 1:1 hexane/EtOAc-100 % EtOAc) to give 4 g (82 %) of the desired compound. ^1H NMR (CDCl_3) δ 1.12 (d, $J = 8.0$ Hz, 18H), 1.25 (t, $J = 7.5$ Hz, 6H), 1.3 (heptet, $J = 8.0$ Hz, 3H), 3.55 (d, $J = 22$

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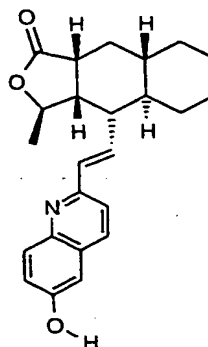
Hz, 2H), 4.08 (q, $J = 7.5$ Hz, 4H), 7.14 (s, 1H), 7.32 (d, $J = 9.5$ Hz, 1H), 7.44 (d, $J = 7.4$ Hz, 1H), 7.90 (d, $J = 7.4$ Hz, 1H), 7.95 (d, $J = 9.5$ Hz, 1H).

Step 9:



A solution of the product of Step 8 (4 g, 8.86 mmol) in THF (20 ml) was cooled to 0°C with stirring. n-Butyllithium (3.5 ml of a 2.5M solution in hexanes, 1 eq) was added dropwise. The resulting solution was stirred for a further 10 min and added to a solution of the product of Step 6 (1.9 g, 8.05 mmol) in THF (20 ml) at 0°C. After 1 hr, NH₄Cl solution (saturated) was added and the mixture extracted with EtOAc. The organic extracts were dried (MgSO₄), concentrated and chromatographed (SiO₂, 1:5-1:3 hexane/EtOAc) to give 2.8 g (65 %) of the title compound. ¹H NMR (CDCl₃) δ 1.12 (d, $J = 8.0$ Hz, 18H), 1.0-1.5 (m, 11H), 1.43 (d, $J = 6.0$ Hz, 3H), 1.73 (br d, $J = 9.5$ Hz, 2H), 1.84 (m, 1H), 1.92 (dd, $J = 9.2, 7.0$ Hz, 1H), 2.40 (m, 2H), 2.71 (q, $J = 6.0$ Hz, 1H), 4.77 (m, 1H), 6.46 (dd, $J = 15.8, 9.6$ Hz, 1H), 6.69 (d, $J = 15.8$ Hz, 1H), 7.13 (s, 1H), 7.31 (d, $J = 6.5$ Hz, 1H), 7.43 (d, $J = 8.5$ Hz, 1H), 7.90 (d, $J = 9.0$ Hz, 1H), 7.94 (d, $J = 8.5$ Hz, 1H).

Step 10:

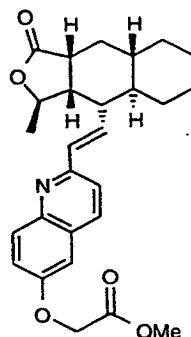


A solution of the product of Step 9 (2.8 g, 5.25 mmol) was stirred in THF (72 ml) at 0°C. Tetrabutylammonium fluoride (5.3 ml of a 1M

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solution in THF, 1 eq) was added dropwise, TLC (5 min) showed complete conversion. NH_4Cl solution (saturated) was added and the mixture extracted with EtOAc. The organic extracts were dried (MgSO_4), concentrated and chromatographed (SiO_2 , 1:2-1:1 hexane/EtOAc) to give 1.96 g (99 %) of the title compound. ^1H NMR (CDCl_3) δ 1.0-1.4 (m, 8H), 1.45 (d, J = 6 Hz, 3H), 1.7-1.9 (m, 3H), 1.97 (m, 1H), 2.43 (m, 2H), 2.72 (q, J = 6.5 Hz, 1H), 4.78 (m, 1H), 6.50 (dd, J = 15.9, 9.5 Hz, 1H), 6.78 (d, J = 15.9 Hz, 1H), 7.17 (s, 1H), 7.34 (d, J = 9.0 Hz, 1H), 7.53 (d, J = 8.5 Hz, 1H), 8.01 (m, 2H); MS m/z 378 (M^+), 332, 264, 236: HRMS calcd for $\text{C}_{24}\text{H}_{28}\text{NO}_3$ (MH^+) 378.2069, found 378.2060. Anal. Calcd for $\text{C}_{24}\text{H}_{27}\text{NO}_3 \cdot \text{HCl} \cdot 0.5\text{H}_2\text{O}$: C, 68.16; H, 6.91; N, 3.31. Found: C, 68.21; H, 7.64; N, 3.36.

Step 11:

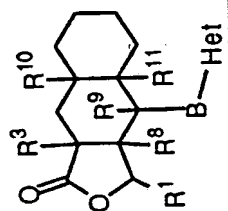


NaH (33 mg of a 60 % dispersion in mineral oil, 0.825 mmol) was added to a solution of the product of Step 10 (75 mg, 0.2 mmol) in DMF (1 ml). After stirring for 10 min, methyl bromoacetate (166 ml, 8 eq) was added and after a further 10 min, TLC indicated complete reaction. NH_4Cl solution (saturated) was added and the mixture extracted with EtOAc. The organic extracts were dried (MgSO_4), concentrated and chromatographed (SiO_2 , 1:4-1:2 hexane/EtOAc) to give 60 mg (68 %) of the title compound. ^1H NMR (CDCl_3) δ 0.8-1.4 (m, 8H), 1.48 (d, J = 6.0 Hz, 3H), 1.79 (br d, J = 9.0 Hz, 2H), 1.89 (d, J = 10.5 Hz, 1H), 1.98 (dd, J = 13.5, 6.0 Hz, 1H), 2.46 (m, 2H), 2.75 (q, J = 6.5 Hz, 1H), 3.89 (s, 3H), 4.82 (br s, 3H), 6.55 (dd, J = 15.7, 9.5 Hz, 1H), 6.58 (d, J = 9.5 Hz, 1H), 7.06 (s, 1H), 7.48 (d, J = 6.5 Hz, 1H), 7.51 (d, J = 8.5 Hz, 1H), 8.03 (t, J = 9.0 Hz, 2H). HRMS calcd for $\text{C}_{27}\text{H}_{32}\text{NO}_5$ (MH^+) 450.2280, found 450.2282.






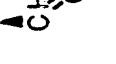






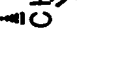






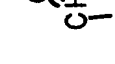
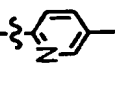





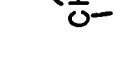






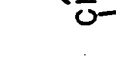
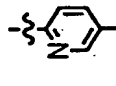





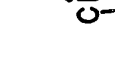







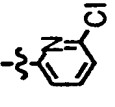
Step 12:

The product of Step 11 (60 mg, 0.136 mmol) was dissolved in 4:1 CH₃OH/water (6.5 ml). LiOH (0.25 ml of a 1 M solution in water, 2 eq) was added and the mixture stirred for 2 hs. Water was added and the mixture extracted with EtOAc. The aqueous layer was acidified and extracted with EtOAc (x3), these extracts were dried (MgSO₄) and concentrated to give 30 mg of the title compound (50 %). ¹H NMR (CD₃OD) δ 0.9-1.5 (m, 8H), 1.42 (d, *J* = 6.0 Hz, 3H), 1.80 (m, 3H), 1.92 (m, 1H), 2.50 (m, 1H), 2.64 (m, 1H), 2.83 (q, *J* = 6.5 Hz, 1H), 4.95 (m, 3H), 6.89 (d, *J* = 15.8 Hz, 1H), 7.16 (dd, *J* = 15.8, 10.0 Hz, 1H), 7.56 (d, *J* = 2.5 Hz, 1H), 7.76 (dd, *J* = 9.5, 2.5 Hz, 1H), 8.08 (d, *J* = 9.5 Hz, 1H), 8.16 (d, *J* = 9.0 Hz, 1H), 8.71 (d, *J* = 9.0 Hz, 1H); MS *m/z* 420 (18), 392 (100), 302 (2), 117 (8); Anal. Calc'd for C₂₆H₂₉NO₅•HCl•2 H₂O: C, 61.47; H, 6.75; N, 2.76. Found: C, 61.02; H, 6.45; N, 2.91.

Using the procedure of Example 1, employing starting materials known in the art or prepared according to procedures similar to those described below in the following examples, the compounds shown in the following Table 1 are prepared, wherein the variables are as defined in the table:








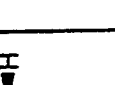






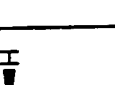
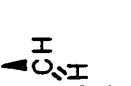





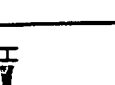
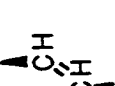





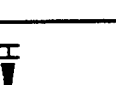
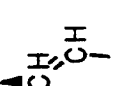





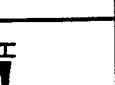
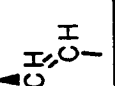






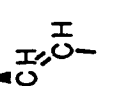
Ex.	R ¹	R ³	R ⁸	R ⁹	R ¹⁰	R ¹¹	B	Het	Physical Data
1A	CH ₃	H	H	H	H	H			HRMS (MH ⁺) found: 340.2281
1B	CH ₃	H	H	H	H	H			MS (CI) m/z = 326 (MH ⁺ , 100%)
1C	CH ₃	H	H	H	H	H			MS (CI) m/z = 326 (MH ⁺ , 100%); [α] _D ²³ = -25.5° (c 0.18, CH ₃ OH)
1D	CH ₃	H	H	H	H	H			MS (CI) m/z = 326 (MH ⁺ , 100%); [α] _D ²³ = +15.8° (c 0.16, CH ₃ OH)
1E	CH ₃	H	H	H	H	H			MS (CI) m/z = 326 (MH ⁺ , 100%)







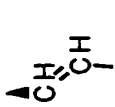
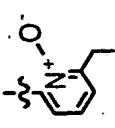






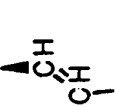
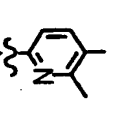






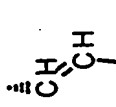
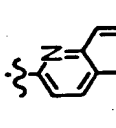






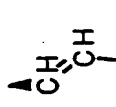
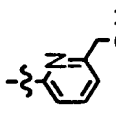






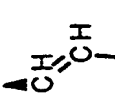
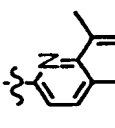






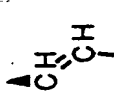
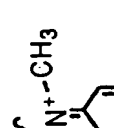






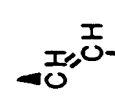
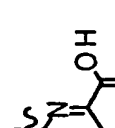
1F	 CH ₃	 H	 H	H	 H	 H	 CH=CH		MS (CI) m/z = 326 (MH ⁺ , 100%); mp = 147-148°C; [α] _D ²⁵ = -32.1° (c 0.52, CH ₃ OH)
1G	 CH ₃	 H	 H	H	 H	 H	 CH=CH		(+)-isomer MS (CI) m/z = 326 (MH ⁺ , 100%)
1H	 CH ₃	 H	 H	H	 H	 H	 CH=CH		HRMS (MH ⁺) found: 326.2118
1I	 CH ₃	 H	 H	H	 H	 H	 CH=CH		HRMS (MH ⁺) found: 326.2115
1J	 CH ₃	 H	 H	H	 H	 H	 CH=CH		HRMS (MH ⁺) found: 326.2118
1K	 CH ₃	 H	 H	H	 H	 H	 CH=CH		HRMS (MH ⁺) found: 326.2115
1L	 CH ₃	 H	 H	H	 H	 H	 CH=CH		MS (CI) m/z = 346 (MH ⁺ , 100%)

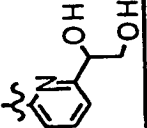
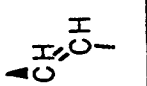




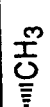
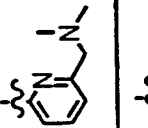
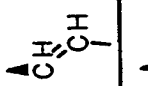




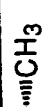
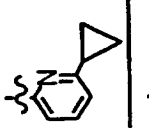
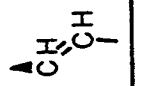




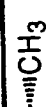
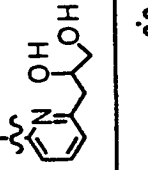
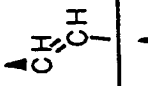




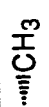
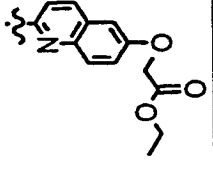
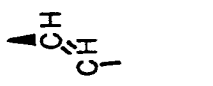




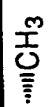
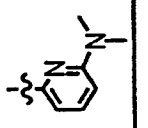
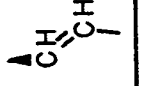





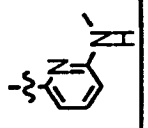
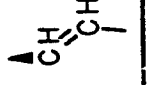





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1M								MS m/z = 328(100), 326(7), 120(7), 107(10)
1N								MS m/z = 328(100), 326(10), 120(6), 107(12)
1O								MS m/z = 328(100), 326(7), 120(7), 107(12)
1P								MS m/z = 328(100), 326(6), 120(9), 107(13)
1Q								MS (Cl) m/z = 312 (MH+, 100%)
1R								MS (Cl) m/z = 368 (MH+, 100%)
1S								MS (Cl) m/z = 326 (MH+, 100%)
1T								MS (Cl) m/z = 395 (MH+, 100%)

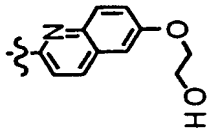
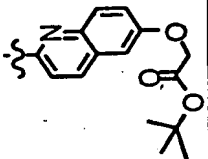
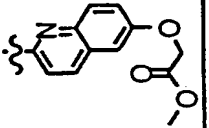
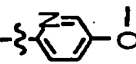
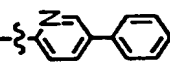
1U									MS (CI) m/z = 410 (MH ⁺ , 100%)
1V									MS (CI) m/z = 417 (MH ⁺ , 100%)
1W									MS (CI) m/z = 362 (MH ⁺ , 100%)
1X									MS (CI) m/z = 340 (MH ⁺ , 100%)
1Y									MS (CI) m/z = 396 (MH ⁺ , 100%)

1AB			H			H				HRMS (MH ⁺) found: 392.2219
1AC			H			H				HRMS (MH ⁺) found: 376.2270
1AD			H			H				HRMS (MH ⁺) found: 392.2219
1AE			H			H				MS (CI) m/z = 354 (MH ⁺ , 100%)
1AF			H			H				MS (CI) m/z = 354 (MH ⁺ , 100%)
1AG			H			H				MS (CI) m/z = 340 (MH ⁺ , 100%); [α] _D ²³ = -25.2° (c 0.24, CH ₃ OH)

1AH									MS (FAB) m/z = 356 (MH ⁺ , 100%)
1AI									HRMS (MH ⁺) found: 340.2285
1AJ									MS (FAB) m/z = 362 (MH ⁺ , 100%)
1AK									MS (FAB) m/z = 342 (MH ⁺ , 100%)
1AL									HRMS (MH ⁺) found: 376.2274
1AM									MS (FAB) m/z = 376 (MH ⁺ , 100%)
1AN									MS (FAB) m/z = 356 (MH ⁺ , 100%)

1AO					H				MS (FAB) m/z = 372 (MH ⁺ , 100%)
1AP					H				MS (FAB) m/z = 369 (MH ⁺ , 100%)
1AQ					H				MS (FAB) m/z = 352 (MH ⁺ , 100%)
1AR					H				MS (FAB) m/z = 386 (MH ⁺ , 100%)
1AS					HH				MS m/z = 464(100)
1AT					H				MS (CI) m/z = 355 (MH ⁺ , 100%)
1AU					H				MS (FAB) m/z = 341 (MH ⁺ , 100%)

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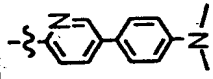
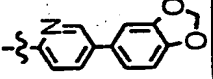
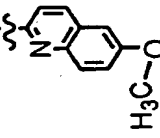
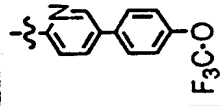
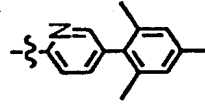
1AV		HRMS (MH ⁺) found: 422.2340
1AW		HRMS (MH ⁺) found: 492.2738
1AX		HRMS (MH ⁺) found: 450.2282
1AY		MS (CI) m/z = 342 (MH ⁺ , 100%)
1AZ		MS (FAB) m/z = 388 (MH ⁺ , 100%)

1BA	CH ₃	H	H	H	H	H	CH ₃	1BA
1BB	CH ₃	H	H	H	H	H	CH ₃	1BB
1BC	CH ₃	H	H	H	H	H	CH ₃	1BC
1BD	CH ₃	H	H	H	H	H	CH ₃	1BD
1BE	CH ₃	H	H	H	H	H	CH ₃	1BE

1BF							MS (FAB) m/z = 396 (MH ⁺ , 100%)
1BG							HRMS (MH ⁺) found: 378.2060
1BH							MS: 406 (M+H ⁺)
1BI							MS (ESI) m/z = 456 (MH ⁺ , 100%)
1BJ							MS (ESI) m/z = 456 (MH ⁺ , 100%)

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1BK								MS (ESI) m/z = 402 (MH ⁺ , 100%)
1BL								MS (CI) m/z = 394 (MH ⁺ , 43%), 188(100%)
1BM								$[\alpha]_D^{25} = -42.88^\circ$ (c 1.0, CH ₃ OH)
1BN								HRMS (MH ⁺) found: 378.2060 $[\alpha]_D^{25} = +39.95^\circ$ (c 1.5, CH ₃ OH)
1BO								MS (FAB) m/z = 438 (MH ⁺ , 100%)

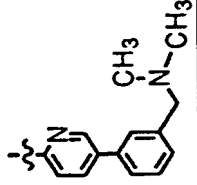
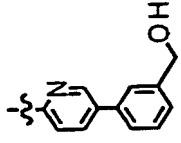
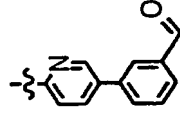
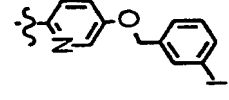
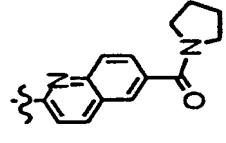
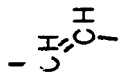
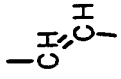
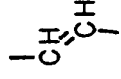
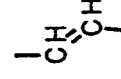
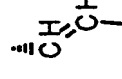






























1BP		MS (CI) m/z = 431 (MH ⁺ , 100%)
1BQ		MS (FAB) m/z = 432 (MH ⁺ , 100%)
1BR		MS (FAB) m/z = 394 (MH ⁺ , 100%)
1BS		MS (FAB) m/z = 472 (MH ⁺ , 100%)
1BT		MS (FAB) m/z = 430 (MH ⁺ , 100%)

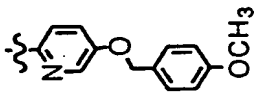
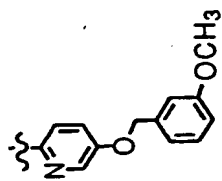
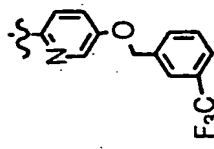
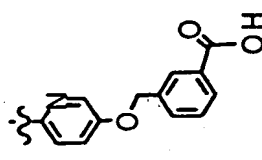
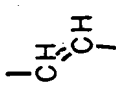
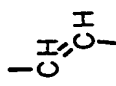
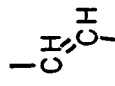
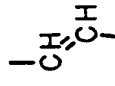
























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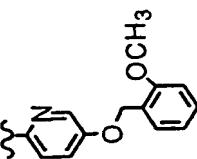





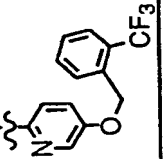





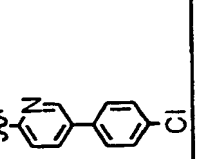





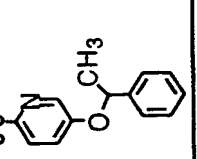





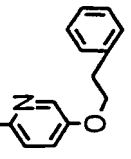
1BU				H			MS m/z = 378(100), 377(15), 154(2), 150(2)
1BV				H			MS m/z = 433(5), 420(18), 393(28), 392(100), 117(8) [α] _D ²³ = +15.7° (c 0.45, CH ₃ OH)
1BW				H			[α] _D ²³ = -14.65° (c 0.45, CH ₃ OH)
1BX				H			HRMS (MH+) found: 392.2225
1BY				H			HRMS (MH+) found: 468.2539
1BZ				H			HRMS (MH+) found: 468.2536






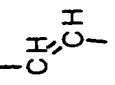
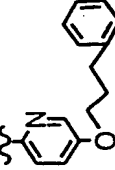





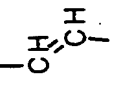
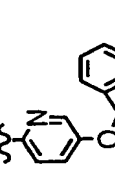





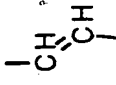
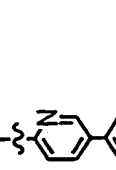





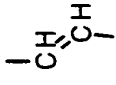
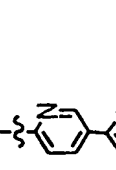





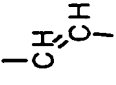
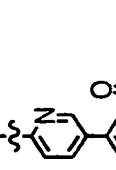





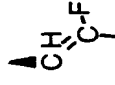
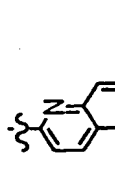
1CA		HRMS (MH ⁺) found: 380.2232
1CB		MS (ESI) m/z = 406 (MH ⁺ , 100%)
1CC		MS (FAB) m/z = 440 (MH ⁺ , 100%)
1CD		MS (FAB) m/z = 4625 (MH ⁺ , 100%)
1CE		MS (FAB) m/z = 419 (MH ⁺ , 100%)
1CF		MS (FAB) m/z = 476 (MH ⁺ , 100%)

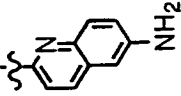
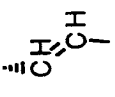






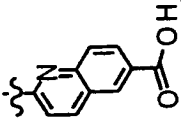
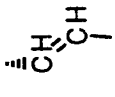






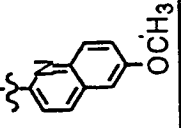
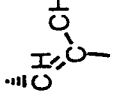






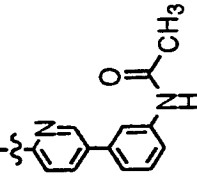
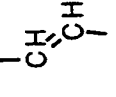






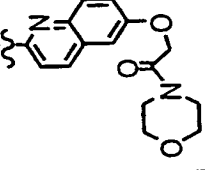
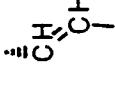






-44-

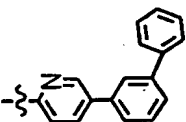
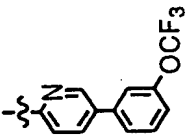
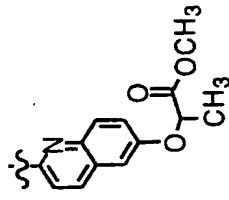
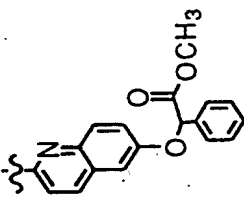
1CG		MS (FAB) m/z = 445 (MH ⁺ , 100%)
1CH		MS (FAB) m/z = 418 (MH ⁺ , 100%)
1CI		MS (FAB) m/z = 416 (MH ⁺ , 100%)
1CJ		MS (FAB) m/z = 544 (MH ⁺ , 100%)
1CK		HRMS (MH ⁺) found: 459.2635
		
		
		
		
		
		
		
		
		
		
		
		
		
		
		
		
		
		
		
		
		
		
		
		
		
		
		
		
		
		
		
		
		
		
		

MS (FAB) m/z = 448 (MH+, 100%)				
				
				
				
				
				
				
				
1CL	1CM	1CN	1CO	

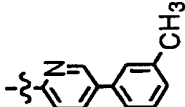
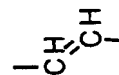






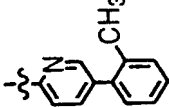
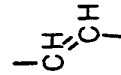






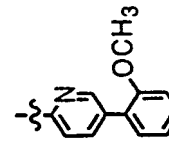
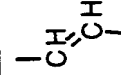






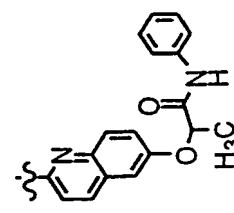
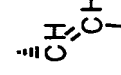






1CP							MS (Cl) m/z = 448 (MH ⁺ , 100%)
1CQ							MS (Cl) m/z = 486 (MH ⁺ , 100%)
1CR							MS (Cl) m/z = 422 (MH ⁺ , 100%)
1CS							MS (Cl) m/z = 432 (MH ⁺ , 100%)
1CT							MS (Cl) m/z = 432 (MH ⁺ , 100%)

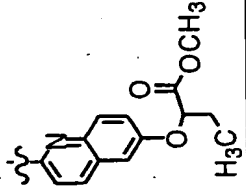
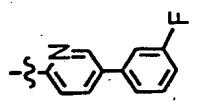
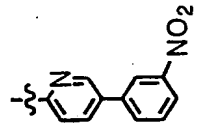
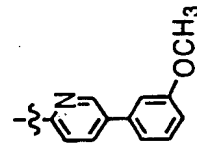
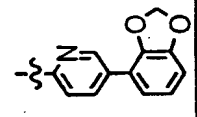
1CU	 CH ₃	 H	 H	 H	 H			MS (FAB) m/z = 446 (MH ⁺ , 100%)
1CV	 CH ₃	 H	 H	 H	 H			MS (CI) m/z = 418 (MH ⁺ , 100%); [α] _D ²⁴ = +26.2° (c 0.46, CH ₃ OH)
1CW	 CH ₃	 H	 H	 H	 H			MS (CI) m/z = 422 (MH ⁺ , 100%); [α] _D ²⁴ = +19.8° (c 0.39, CH ₃ OH)
1CX	 CH ₃	 H	 H	 H	 H			MS (CI) m/z = 394 (MH ⁺ , 100%); [α] _D ²⁴ = +23.8° (c 0.20, CH ₃ OH)
1CY	 CH ₃	 H	 H	 H	 H			MS (FAB) m/z = 432 (MH ⁺ , 100%)
1CZ	 CH ₃	 H	 H	 H	 H			MS: 396 (M+H ⁺)

1DA									HRMS (MH+) found: 377.2224
1DB									MS m/z = 406(100), 241(25), 225(51), 194(56), 168(53)
1DC									HRMS (MH+) found: 406.2381
1DD									MS (Cl) m/z = 445 (MH+, 100%)
1DE									HRMS (MH+) found: 505.2698







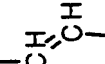
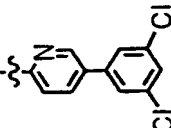





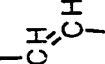
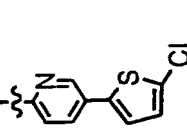





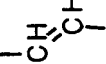
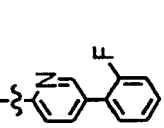





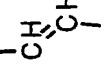
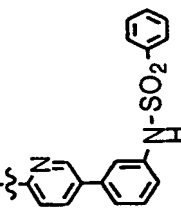





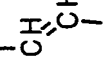
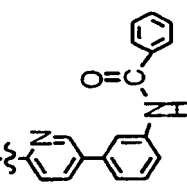
1DF		MS (FAB) m/z = 464 (MH ⁺ , 100%)
1DG		MS (FAB) m/z = 472 (MH ⁺ , 100%)
1DH		HRMS (MH ⁺) found: 464.2431
1DI		HRMS (MH ⁺) found: 526.2592

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



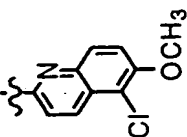








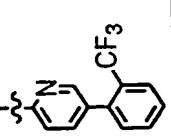








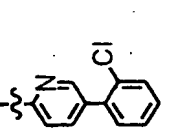








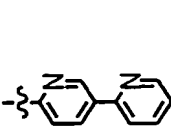








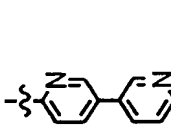
1DJ									MS(ESI) m/z = 402 (MH+, 100%); [α] _D ²² = +23.2° (c 0.55, CH ₃ OH)
1DK									MS(ESI) m/z = 402 (MH+, 100%); [α] _D ²² = +18.1° (c 0.42, CH ₃ OH)
1DL									MS(FAB) m/z = 418 (MH+, 100%); [α] _D ²³ = +19.5° (c 0.86, CH ₃ OH)
1DM									HRMS (MH+) found: 525.2746

1DN		HRMS (MH+) found: 478.2584
1DO		MS (ESI) m/z = 418 (MH+, 100%) [α] _D ²³ = +23.9° (c 0.38, CH ₃ OH)
1DP		MS (ESI) m/z = 433 (MH+, 100%) [α] _D ²² = +22.3° (c 0.44, CH ₃ OH)
1DQ		MS (ESI) m/z = 418 (MH+, 100%)
1DR		MS (CI) m/z = 432 (MH+, 100%)







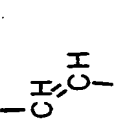
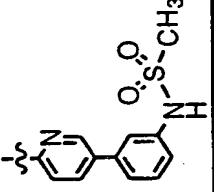






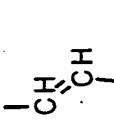
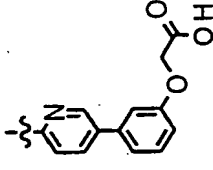






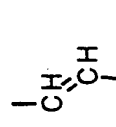
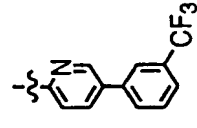






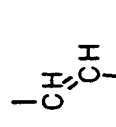
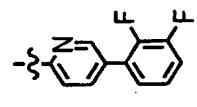






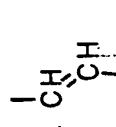
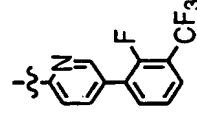
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1DS						 		MS (ESI) m/z = 456 (MH ⁺ , 100%) [α] _D ²² = +20.3° (c 0.30, CH ₃ OH)
1DT								MS (ESI) m/z = 428 (MH ⁺ , 100%) [α] _D ²⁴ = +25.4° (c 0.26, CH ₃ OH)
1DU								MS (CI) m/z = 406 (MH ⁺ , 100%) [α] _D ²⁴ = +17.2° (c 0.50, CH ₃ OH)
1DV								MS (FAB) m/z = 543 (MH ⁺ , 100%)
1DW								MS (FAB) m/z = 507 (MH ⁺ , 100%)

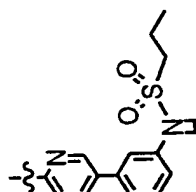
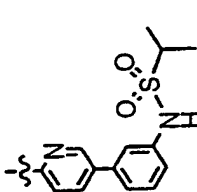
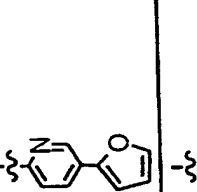
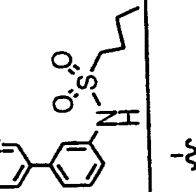
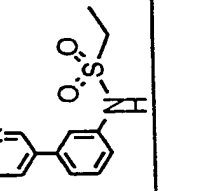
-53-

1DX							 		MS (CI) m/z = 426 (MH ⁺ , 100%)
1DY							 		MS (CI) m/z = 456 (MH ⁺ , 100%)
1DZ							 		MS (CI) m/z = 422 (MH ⁺ , 95%) [α] _D ²³ = +15.1° (c 0.35, CH ₃ OH)
1EA							 		MS (CI) m/z = 389 (MH ⁺ , 100%)
1EB							 		MS (FAB) m/z = 389 (MH ⁺ , 100%)







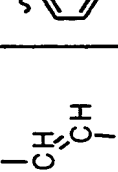
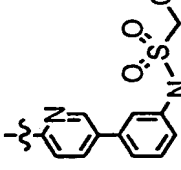






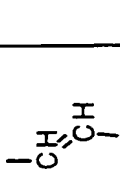
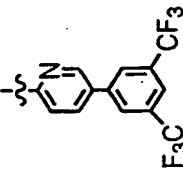






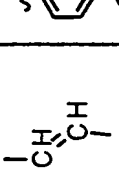
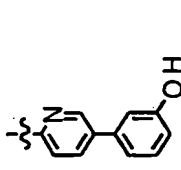






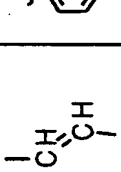
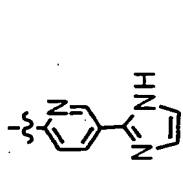






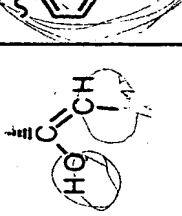
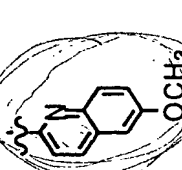
1EC							MS (FAB) m/z = 388 (MH+, 100%)
1ED							MS (CI) m/z = 430 (MH+, 100%)
1FE							MS (CI) m/z = 430 (MH+, 100%)
1EF							MS (CI) m/z = 402 (MH+, 100%) [α] _D ²⁴ = +15.7° (c 0.33, CHCl ₃)
1EG							MS (CI) m/z = 476 (MH+, 100%)

1EH	 CH ₃	 H	 H	 H	 H	 H			MS (FAB) m/z = 481 (MH ⁺ , 100%)
1EI	 CH ₃	 H	 H	 H	 H	 H			MS (ESI) m/z = 462 (MH ⁺ , 100%)
1EJ	 CH ₃	 H	 H	 H	 H	 H			MS (ESI) m/z = 456 (MH ⁺ , 100%) [α] _D ²⁰ = +40.0° (c 0.39, CHCl ₃)
1EK	 CH ₃	 H	 H	 H	 H	 H			MS (ESI) m/z = 424 (MH ⁺ , 100%) [α] _D ²² = +25.5° (c 0.26, CH ₃ OH)
1EL	 CH ₃	 H	 H	 H	 H	 H			MS (ESI) m/z = 474 (MH ⁺ , 100%) [α] _D ²² = +22.4° (c 0.50, CH ₃ OH)

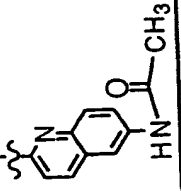
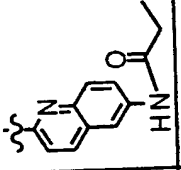
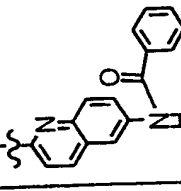
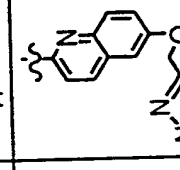
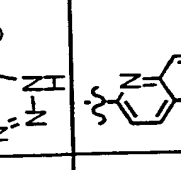
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1EM	CH ₃	H	H	H	H		MS (FAB) m/z = 509 (MH ⁺ , 100%)
1EN	CH ₃	H	H	H	H		MS (FAB) m/z = 509 (MH ⁺ , 100%)
1EO	CH ₃	H	H	H	H		MS (ESI) m/z = 378 (MH ⁺ , 100%)
1EP	CH ₃	H	H	H	H		MS (ESI) m/z = 523 (MH ⁺ , 100%)
1EQ	CH ₃	H	H	H	H		MS (ESI) m/z = 495 (MH ⁺ , 100%)

B

1ER									MS (ESI) m/z = 549 (MH ⁺ , 100%)
1ES									MS (FAB) m/z = 524 (MH ⁺ , 100%)
1ET									MS (CI) m/z = 404 (MH ⁺ , 100%) [α] _D ²³ = +32.6° (c 0.27, CH ₃ OH)
1EU									MS (ESI) m/z = 378 (MH ⁺ , 100%)
1EV									HRMS (MH ⁺) found 408,2174

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1EW		HRMS (MH+) found 419.2331
1EX		HRMS (MH+) found 433.2489
1EY		HRMS (MH+) found 481.2495
1EZ		HRMS (MH+) found 460.2335
1FA		HRMS (MH+) found 478.1611

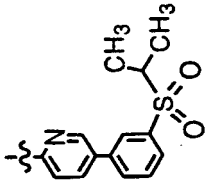
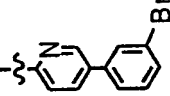
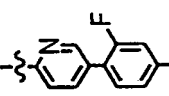
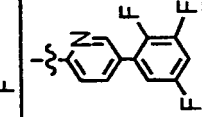
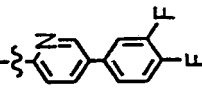
1FB				H				HRMS (MH+) found 444.1998
1FC				H				HRMS (MH+) found 412.2277
1FD				H				MS: 323 (M+H)
1FE				H				MS: 309 (M+1)
1FF				H				HRMS (MH+) found 376.2266
1FG				H				MS (FAB) m/z = 343 (MH+, 100%)

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

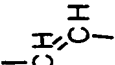
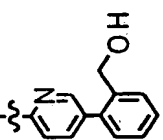





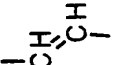
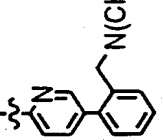





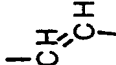
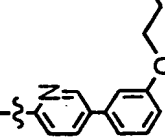





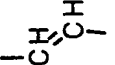
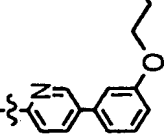





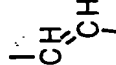
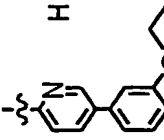
1FH	\blacktriangleleft CH ₃	\blacktriangledown H	\blacktriangledown H	\blacktriangledown H	\blacktriangledown H	$\text{---}\text{CH}=\text{CH}\text{---}$		MS (CI) m/z = 424 (MH ⁺ , 100%)
1FI	\blacktriangleleft CH ₃	\blacktriangledown H	\blacktriangledown H	\blacktriangledown H	\blacktriangledown H	$\text{---}\text{CH}=\text{CH}\text{---}$		MS (ESI) m/z = 456 (MH ⁺ , 100%)
1FJ	\blacktriangleleft CH ₃	\blacktriangledown H	\blacktriangledown H	\blacktriangledown H	\blacktriangledown H	$\text{---}\text{CH}=\text{CH}\text{---}$		MS (CI) m/z = 460 (MH ⁺ , 100%)
1FK	\blacktriangleleft CH ₃	\blacktriangledown H	\blacktriangledown H	\blacktriangledown H	\blacktriangledown H	$\text{---}\text{CH}=\text{CH}\text{---}$		MS (ESI) m/z = 474 (MH ⁺ , 100%)
1FL	\blacktriangleleft CH ₃	\blacktriangledown H	\blacktriangledown H	\blacktriangledown H	\blacktriangledown H	$\text{---}\text{CH}=\text{CH}\text{---}$		MS (ESI) m/z = 413 (MH ⁺ , 100%)

1FM								MS (FAB) m/z = 460 (MH ⁺ , 100%)
1FN								MS (ESI) m/z = 424 (MH ⁺ , 100%)
1FO								MS (ESI) m/z = 472 (MH ⁺ , 100%)
1FP								MS (ESI) m/z = 424 (MH ⁺ , 100%)
1FQ								MS (ESI) m/z = 467 (MH ⁺ , 100%)

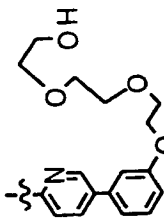
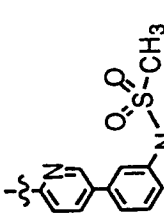
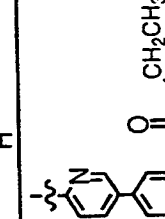
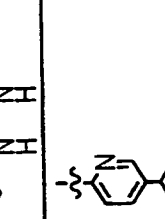
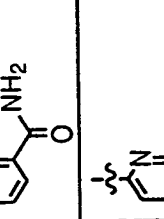
-62-

1FR		MS (ESI) m/z = 494 (MH ⁺ , 100%)
1FS		MS (FAB) m/z = 466 (MH ⁺ , 100%)
1FT		MS (FAB) m/z = 424 (MH ⁺ , 100%)
1FU		MS (FAB) m/z = 442 (MH ⁺ , 100%)
1FV		MS (CI) m/z = 424 (MH ⁺ , 100%)

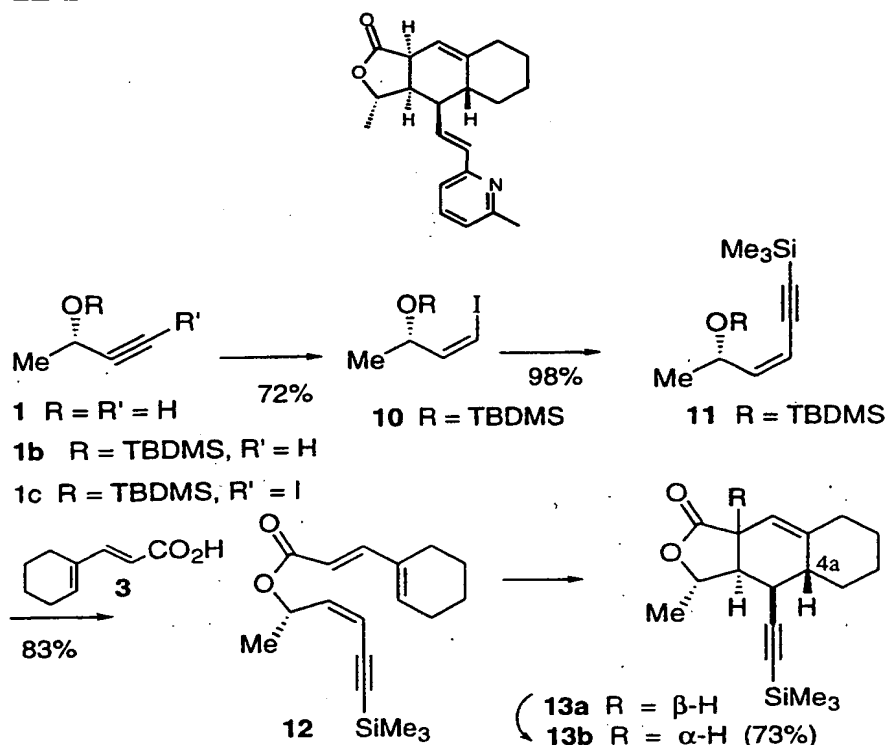
-63-

1FW								MS (ESI) m/z = 418 (MH ⁺ , 100%)
1FX								MS (ESI) m/z = 445 (MH ⁺ , 100%)
1FY								MS (ESI) m/z = 448 (MH ⁺ , 100%)
1FZ								MS (ESI) m/z = 462 (MH ⁺ , 100%)
1GA								MS (ESI) m/z = 492 (MH ⁺ , 100%)

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1GB		MS (ESI) m/z = 536 (MH ⁺ , 100%)
1GC		MS (ESI) m/z = 481 (MH ⁺ , 100%)
1GD		MS (ESI) m/z = 474 (MH ⁺ , 100%)
1GE		MS (FAB) m/z = 431 (MH ⁺ , 100%)
1GF		MS (FAB) m/z = 430 (MH ⁺ , 100%)

EXAMPLE 2



- Step 1:** To a solution of compound **1b** (4.532 g, 24.6 mmol) in THF (15 ml), cooled to 0 °C under argon, was added a solution of *n*-BuLi (1.6 M in cyclohexane, 17 ml, 27 mmol). After stirring at 0 °C for 40 min, a solution of I₂ (6.24 g, 24.6 mmol) in THF (10 ml) was added and the reaction mixture was stirred for an additional 15 min. The reaction was quenched by addition of water (25 ml) and diluted with hexane (50 ml).
- The aqueous phase was extracted with hexane (3 x 50 ml). The combined organic phase was washed with 5% sodium thiosulfate solution (2 x 50 ml), dried over MgSO₄ and evaporated *in vacuo* to give the acetylenic iodide **1c** as an orange oil (7.281 g, 95%).
- [α]_D²³ -48.8 (*c* 1.23, CHCl₃); IR (CH₂Cl₂) 2200 cm⁻¹; ¹H NMR (400 MHz, CDCl₃) δ 0.11 (s, 3 H), 0.12 (s, 3 H), 0.90 (s, 9 H, C(CH₃)₃), 1.40 (d, *J*=6.5 Hz, 3 H, CH₃), 4.63 (q, *J*=6.4 Hz, 1 H, CH(OTBS)); ¹³C NMR (100 MHz, CDCl₃) δ -4.65, -0.31, 18.21, 25.35, 25.76, 60.50, 96.98.

- Step 2:** To a solution of cyclohexene (6.8 ml, 67 mmol) in anhydrous pentane (50 ml) stirred at 0 °C under argon was added borane-methyl sulfide complex (2 M in THF, 16.7 ml, 33.4 mmol). The solution was warmed to room temperature and stirred for 1 h to give a

cloudy suspension to which was added the acetylenic iodide **1c** (8.446 g, 27.2 mmol). The resulting clear solution was stirred at room temperature for 80 min, and glacial acetic acid (5 ml, 87.3 mmol) was added. After stirring the reaction mixture for 20 min, ethanolamine (5.2 ml, 86.2 mmol) was added and stirring was continued for an additional 15 min. The mixture was diluted with EtOAc (300 ml), washed with water (2 x 100 ml), and brine (100 ml). The organic phase was dried over anhydrous MgSO₄ and concentrated *in vacuo* to yield the crude product as a yellow oil. Purification by column chromatography on silica gel (hexane) gave *cis*-vinyl iodide **10** as a colorless oil (7.167 g, 84%).
[α]_D²³ +68.1 (*c* 0.79, CHCl₃); IR (KBr pellet) 1610 cm⁻¹; ¹H NMR (400 MHz, CDCl₃) δ 0.13 (s, 3 H), 0.16 (s, 3 H), 0.95 (s, 9 H, C(CH₃)₃), 1.27 (d, *J*=6.4, 3 H, CH₃), 4.56 (dq, *J*=6.4, 6.2 Hz, 1 H, OCH), 6.18 (d, *J*=7.6 Hz, 1 H, HC=CH_I), 6.28 (dd, *J*=7.6, 7.6 Hz, 1 H, CH=CH_I); ¹³C NMR (100 MHz, CDCl₃) δ -5.15, -4.95, 17.80, 22.37, 25.29, 71.63, 78.25, 145.09.

Step 3: To a solution of PdCl₂(PhCN)₂ (58.1 mg, 0.15 mmol) and CuI (58.8 mg, 0.31 mmol) in piperidine (3 ml) was added a solution of the *cis*-vinyl iodide **10** (303 mg, 0.97 mmol) in anhydrous THF (3 ml). This was followed by addition of (trimethylsilyl)acetylene (0.35 ml, 2.48 mmol) which was accompanied by a color change from dark green to pale green and then to black over 5 min. The solution was stirred at room temperature under argon for 18 h. The solvents were removed *in vacuo* and the mixture was purified by flash chromatography on silica gel (hexane, followed by 5% EtOAc in hexane) to give the product **11** as a yellow oil (267 mg, 98%).

[α]_D²⁵ +128.7 (*c* 0.745, CHCl₃); IR (CH₂Cl₂) 2151, 1252 cm⁻¹; ¹H NMR (400 MHz, CDCl₃) δ 0.12 (s, 3 H), 0.15 (s, 3 H), 0.25 (s, 9 H, Si(CH₃)₃), 0.95 (s, 9 H, C(CH₃)₃), 1.29 (d, *J*=6.2 Hz, 3 H, CH₃), 4.89 (dq, *J*=8.5, 6.3 Hz, 1 H, OCH), 5.46 (d, *J*=11.0 Hz, 1 H, CH=C), 5.97 (dd, *J*=8.5, 11.0 Hz, 1 H, C=CH); ¹³C NMR (100 MHz, CDCl₃) δ -4.88, -4.58, 1.08, 18.21, 23.61, 25.91, 67.03, 99.53, 101.13, 107.40, 148.86.; MS (CI/CH₄) 283 (MH⁺), 267, 225.

Step 4: To a solution of the protected enyne **11** (1.744 g, 10.38 mmol) in CH₃OH (30 ml) was added TFA (0.6 ml). The reaction mixture was stirred at rt for 2 h. The solvent was removed *in vacuo* and the residue was diluted with Et₂O (40 ml) and water (40 ml). The aqueous

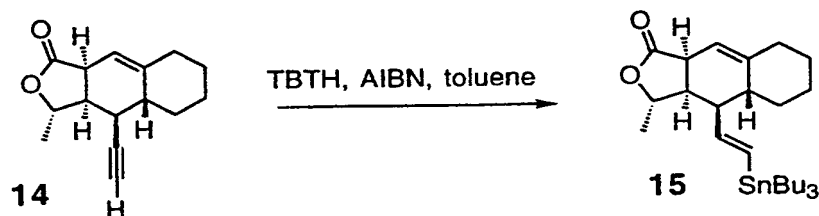
phase was extracted with Et₂O (3 x 40 ml), the combined organic phase was washed with brine (50 ml), dried over MgSO₄, and concentrated. The final trace of solvent was removed under high vacuum.

To a solution of the deprotected enyne from above in anhydrous CH₂Cl₂ (30 ml) was added 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (4.412 g, 23.01 mmol), dimethylaminopyridine (DMAP) (2.836 g, 23.2 mmol), TEMPO (1 mg), and dienoic acid **3** (2.414 g, 15.9 mmol). The reaction mixture was stirred at room temperature under argon for 18 h. The solvents were removed and the mixture was diluted with EtOAc (300 ml). The organic phase was washed with water (150 ml), 0.5 N HCl solution (2 x 100 ml), and brine (100 ml) and dried over anhydrous MgSO₄. Evaporation under reduced pressure yielded the ester **12** as a brown oil (2.601 g, 83%). An analytical sample was prepared by further chromatography on silica gel (5% EtOAc in hexane).
[α]_D²⁵ +190.7 (c 1.04, CHCl₃); IR (CH₂Cl₂) 2151, 1715 cm⁻¹; ¹H NMR (400 MHz, CDCl₃) δ 0.20 (s, 9 H, C(CH₃)₃), 1.40 (d, J=6.4 Hz, 3 H, CH₃), 1.58-1.66 (m, 2 H, CH₂), 1.66-1.73 (m, 2 H, CH₂), 2.10-2.18 (m, 2 H, CH₂), 2.23-2.30 (m, 2 H, CH₂), 5.57 (dd, J=11.0, 1.1 Hz, 1 H, C=CH), 5.76 (d, J=15.6 Hz, 1 H, CH=C), 5.86 (dq, J=6.44, 7.56 Hz, 1 H, CH=C), 5.97 (dd, J=7.8, 11.0 Hz, 1 H, CH=C), 6.22 (t (broad), J=4.0 Hz, CH=C), 7.31 (s, 1 H, CH=C); ¹³C NMR (100 MHz, CDCl₃) δ -0.203, 19.81, 21.98, 22.01, 24.07, 26.40, 69.02, 100.20, 101.51, 110.35, 114.55, 134.85, 138.77, 143.55, 148.21, 166.65; HRMS (FAB): Calcd for C₁₈H₂₇O₂Si (M⁺) m/e 302.1702, found m/e 302.1695.

Step 5: To a solution of intermediate **12** (2.125 g, 7.03 mmol) in anhydrous, degassed toluene (25 ml) was added TEMPO (1 mg). The solution was heated in a sealed tube at 185 °C for 2.5 h. The reaction mixture was cooled to room temperature, added DBU (1 ml) and stirred for 30 min. The mixture was diluted with EtOAc (300 ml) and washed with water (100 ml), 0.5 N HCl solution (2 x 100 ml), and brine (100 ml). The organic phase was dried over anhydrous MgSO₄, filtered and evaporated to give the crude product as a yellow oil (2.290 g). Purification by chromatography on silica gel (8% EtOAc in hexane) gave the tricyclic derivative **13b** as a pale yellow oil (1.541 g, 73%).

[α]_D²¹ +115.6 (c 1.01, CHCl₃); IR (CH₂Cl₂) 2170, 1768 cm⁻¹; ¹H NMR (400 MHz, CDCl₃) δ 0.21 (s, 9 H, Si(CH₃)₃), 0.98 (dddd, J =12.0, 10.5, 10.5, 3.5 Hz, 1 H, C₅H_{ax}), 0.92-1.04 (m, 1 H, CH), 1.23-1.36 (m, 1 H), 1.40-1.54 (m, 1 H), 1.66 (d, J =6.1 Hz, 3 H, CH₃), 1.78-1.94 (m, 2 H), 1.96-2.15 (m, 2 H), 2.31-2.44 (m, 2 H), 2.54-2.68 (m, 2 H), 3.23-3.29 (m, 1 H, C(O)CH), 4.52-4.62 (m, 1 H, OCH(CH₃)), 5.35 (d, J =2.2 Hz, 1 H, C=CH); ¹³C NMR (100 MHz, CDCl₃) δ -0.10, 21.29, 25.80, 26.79, 32.94, 33.18, 34.87, 38.16, 43.49, 44.73, 77.67, 88.22, 107.05, 113.12, 142.13, 175.75; HRMS (FAB) Calcd for C₁₈H₂₇O₂Si (MH⁺) m/e 303.1780, found m/e 303.1775.

Step 6:

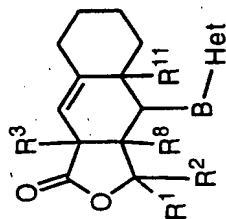


Alkyne **14** (1 g), generated by desilylation of **13b** using K₂CO₃ in CH₃OH, was dissolved in toluene (20 ml) in the presence of tributyltin hydride (1.75 ml) and AIBN (100 mg), and the mixture heated at 120 °C for 2 h. The solution was poured onto a column of silica gel, and the desired product eluted with EtOAc-hexane (5:95). ¹H-NMR (CDCl₃): δ 0.8-0.9 (m, 9H); 1.2-1.6 (m); 3.2 (m, 1H); 4.5 (m, 1H); 5.3 (s, 1H); 5.75 (dd, J = 8.3, 18 Hz, 1H); 6.05 (d, J = 18 Hz, 1H).





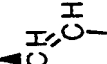











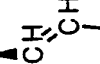
























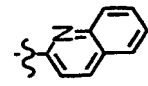




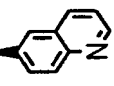
Step 7:

A solution of **15** (224 mg), 6-bromopicoline (356 mg), and Pd(PPh₃)₄ (50 mg) in toluene (4 ml) was heated overnight in pressure tube at 120 °C. The resulting solution was poured onto a column of silica gel and the title compound eluted with EtOAc-hexane (5:95 to 10:90) mixtures. ¹H-NMR (CDCl₃): δ 1.35 (d, 6 Hz, 3H); 2.46 (s, 3H); 3.2 (m, 1H); 4.5 (m, 1H); 5.3 (s, 1H); 6.5 (m, 2H); 6.9 (d, 1H); 7.1 (d, 1H); 7.5 (t, 1H).

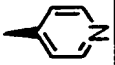
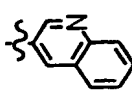
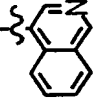
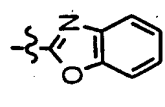
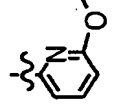
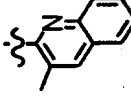
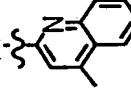
Using a similar procedure, compounds of the following structure are prepared, wherein the variables are as defined in the following Table 2:


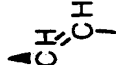
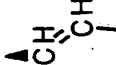
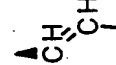
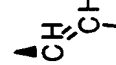
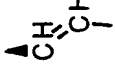
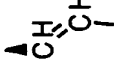






























Ex.	R ¹	R ²	R ³	R ⁸	R ¹¹	B	Het	Physical Data
2A	CH ₃	H	H	H	H			MS: 378 (M+H ⁺)
2B	CH ₃	H	H	H	H			MS: 342 (M+H ⁺)
2C	CH ₃	H	H	H	H			MS: 344 (M+H ⁺)
2D	CH ₃	H	H	H	H	-		MS: 334 (M+H ⁺)
2E	CH ₃	H	H	H	H			MS: 324 (M+H ⁺)

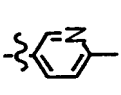
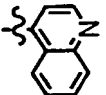

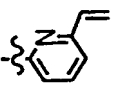
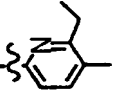
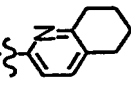
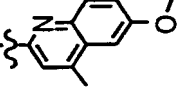
2F		H						MS: 310 (M+H ⁺)
2G		H						MS: 310 (M+H ⁺)
2H		H						MS: 311 (M+H ⁺)
2I		H						MS: 316 (M+H ⁺)
2J		H						MS: 326 (M+H ⁺)
2K		H						MS: 311 (M+H ⁺)
2L		H						MS: 360 (M+H ⁺)
2M		H				-		MS: 334 (M+H ⁺)












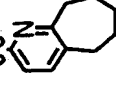





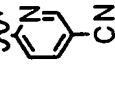





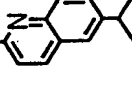





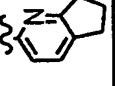





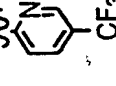





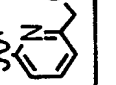
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2N		MS: 284 (M+H ⁺)
2O		MS: 360 (M+H ⁺)
2P		MS: 360 (M+H ⁺)
2Q		MS: 350 (M+H ⁺)
2R		MS: 340 (M+H ⁺)
2S		MS: 374 (M+H ⁺)
2T		MS: 374 (M+H ⁺)

						
						
						
						
H	H	H	H	H	H	H
						

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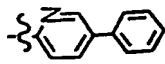




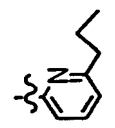




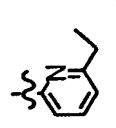




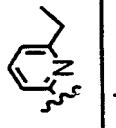




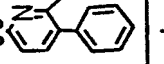




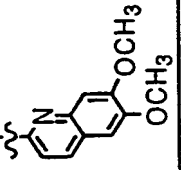




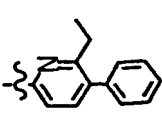
2U	$\cdots\text{CH}_3$	H	$\cdots\text{H}$	$\cdots\text{H}$	H	$\text{CH}_2=\text{CH}-$		MS: 324 (M+H ⁺)
2V	$\cdots\text{CH}_3$	H	$\cdots\text{H}$	$\cdots\text{H}$	H	$\text{CH}_2=\text{CH}-$		MS: 360 (M+H ⁺)
2W	$\cdots\text{CH}_3$	H	$\cdots\text{H}$	$\cdots\text{H}$	H	$\text{CH}_2=\text{CH}-$		MS: 364 (M+H ⁺)
2X	$\cdots\text{CH}_3$	H	$\cdots\text{H}$	$\cdots\text{H}$	H	$\text{CH}_2=\text{CH}-$		MS: 336 (M+H ⁺)
2Y	$\cdots\text{CH}_3$	H	$\cdots\text{H}$	$\cdots\text{H}$	H	$\text{CH}_2=\text{CH}-$		MS: 352 (M+H ⁺)
2Z	$\cdots\text{CH}_3$	H	$\cdots\text{H}$	$\cdots\text{H}$	H	$\text{CH}_2=\text{CH}-$		MS: 364 (M+H ⁺)
2AA	$\cdots\text{CH}_3$	H	$\cdots\text{H}$	$\cdots\text{H}$	H	$\text{CH}_2=\text{CH}-$		MS: 404 (M+H ⁺)

2AB		H						MS: 338 (M+H ⁺)
2AC		H						MS: 378 (M+H ⁺)
2AD		H						MS: 335 (M+H ⁺)
2AE		H						MS: 402 (M+H ⁺)
2AF		H						MS: 350 (M+H ⁺)
2AG		H						MS: 378 (M+H ⁺)
2AH		H						MS: 354 (M+H ⁺)






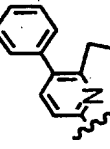




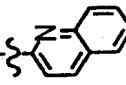
















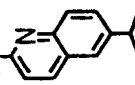





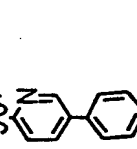





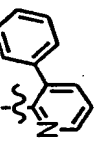
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2AI	$\cdots\text{CH}_3$	H	$\cdots\text{H}$	$\cdots\text{H}$	$\text{H} \blacktriangledown$	$\text{CH}=\text{CH}$		MS: 325 (M+H ⁺)
2AJ	$\cdots\text{CH}_3$	H	$\cdots\text{H}$	$\cdots\text{H}$	$\text{H} \blacktriangledown$	$\text{CH}=\text{CH}$		MS: 360 (M+H ⁺)
2AK	$\cdots\text{CH}_3$	H	$\cdots\text{H}$	$\cdots\text{H}$	$\text{H} \blacktriangledown$	$\text{C}=\text{C}$		MS: 360 (M+H ⁺)
2AL	CH_3CH_2	H	$\cdots\text{H}$	$\cdots\text{H}$	$\text{H} \blacktriangledown$	$\text{CH}=\text{CH}$		MS: 374 (M+H ⁺)
2AM	$\cdots\text{CH}_3$	H	$\cdots\text{H}$	$\cdots\text{H}$	$\text{H} \blacktriangledown$	$\text{CH}=\text{CH}$		MS: 361 (M+H ⁺)
2AN	$\cdots\text{CH}_3$	H	$\cdots\text{H}$	$\cdots\text{H}$	$\text{H} \blacktriangledown$	$\text{CH}=\text{CH}$		MS: 368 (M+H ⁺)
2AO	$-\text{CH}_3$	$-\text{CH}_3$	$\cdots\text{H}$	$\cdots\text{H}$	$\text{H} \blacktriangledown$	$\text{CH}=\text{CH}$		MS: 374 (M+H ⁺)

2AP		MS: 362 (M+H ⁺)
2AQ		MS: 422 (M+H ⁺)
2AR		HRMS found: 422.1944
2AS		MS: 418 (M+H ⁺)
2AT		HRMS found: 368.2224
2AU		HRMS (M+H ⁺) found: 404.1858
2AV		HRMS (M+H ⁺) found: 356.1685
2AW		HRMS (M+H ⁺) found: 356.1685

2AX		H		H		H			HRMS (M+H ⁺) found: 386.2115
2AY		H		H		H			HRMS (M+H ⁺) found: 352.2274
2AZ		H		H		H			HRMS (M+H ⁺) found: 338.2113
2BA		H		H		H			HRMS (M+H ⁺) found: 338.2110
2BB		H		H		H			HRMS (M+H ⁺) found: 400.2273
2BC		H		H		H			HRMS (M+H ⁺) found: 420.2181
2BD		H		H		H			HRMS (M+H ⁺) found: 414.2432

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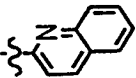
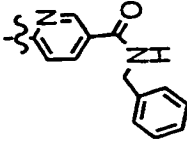
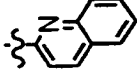
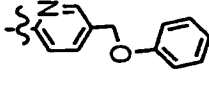
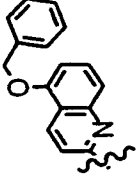
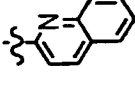
2BE		H	H						HRMS (M+H ⁺) found: 414.2432
2BF	H	H						HRMS (M+H ⁺) found: 346.1811	
2BG	H	H						HRMS (M+H ⁺) found: 310.1808	
2BH	 CH ₃ CH ₂	H						HRMS (M+H ⁺) found: 338.2127	
2BI	 CH ₃	H						HRMS (M+H ⁺) found: 416.2593	
2BJ	 CH ₃	H						HRMS (M+H ⁺) found: 386.2115	
2BK	 CH ₃	H						HRMS (M+H ⁺) found: 386.2115	

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2BL		HRMS (M+H ⁺) found: 380.2594
2BM		HRMS (M+H ⁺) found: 434.2331
2BN		HRMS (M+H ⁺) found: 404.1867
2BO		HRMS (M+H ⁺) found: 404.1871
2BP		HRMS (M+H ⁺) found: 410.2126
2BQ		HRMS (M+H ⁺) found: 410.2122

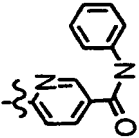
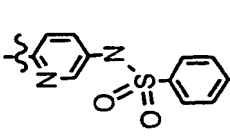
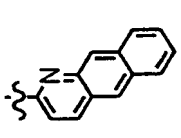
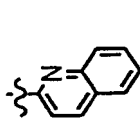
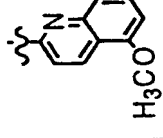
-79-

2BR		HRMS (M+H ⁺) found: 400.2269
2BS		HRMS (M+H ⁺) found: 436.2271
2BT		MS: 388 (M+H ⁺)
2BU		HRMS (M+H ⁺) found: 411.2072
2BV		HRMS (M+H ⁺) found: 390.2064
2BW		HRMS (M+H ⁺) found: 400.2293

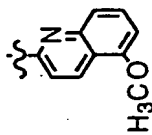
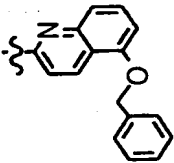

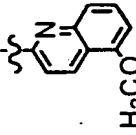
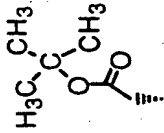
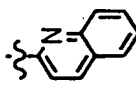
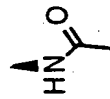
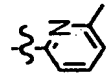
2BX		HRMS (M+H ⁺) found: 374.2107
2BY		HRMS (M+H ⁺) found: 443.2336
2BZ		HRMS (M+H ⁺) found: 450.2445
2CA		MS: 416 (M+H ⁺)
2CB		HRMS (M+H ⁺) found: 466.2373
2CC		HRMS (M+H ⁺) found: 418.2022

-81-

2CD		H						HRMS (M+H ⁺) found: 430.2375
2CE		H						MS: 430 (M+H ⁺)
2CF		H						MS: 429 (M+H ⁺)
2CG		H						MS: 432 (M+H ⁺)
2CH		H						MS: 435 (M+H ⁺)
2CI		H						MS: 416 (M+H ⁺)

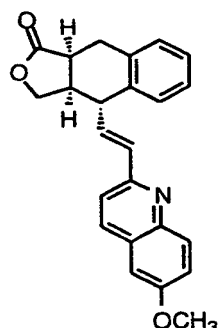
2CJ		MS: 429 (M+H ⁺)
2CK		MS: 465 (M+H ⁺)
2CL		HRMS (M+H ⁺) found: 410.2130
2CM		HRMS (M+H ⁺) found: 404.2226
2CN		HRMS (M+H ⁺) found: 390.2071

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2CO	CH_3	H	CH_3	H	CH_3	H	$\text{CH}=\text{CH}$		HRMS (M+H ⁺) found: 404.2231
2CP	CH_3	H	H	H	H	H	$\text{CH}=\text{CH}$		HRMS (M+H ⁺) found: 466.2377
2CQ	CH_3	H	H	H	H	H			HRMS (M+H ⁺) found: 392.2223
2CR	CH_3	H		H	H	H	$\text{CH}=\text{CH}$		MS: 474 (M+H ⁺)
2CS	CH_3	H	H	H	H	H			MS (FAB): 343 (MH ⁺ , 100%)

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EXAMPLE 3



The compound numbers used in the following steps correspond to those shown in reaction Scheme 3, above.

- 5 Step 1: A solution of THP-ether **18** (2.8 g, 20 mmol) in dry THF (100 ml) was cooled to -78 °C and *n*-BuLi (25 mmol, 10 ml, 2.5N in hexanes) was added dropwise. After 15 min at that temperature, benzyl chloroformate (3.75 ml, 25 mmol, 95% pure) was added dropwise. The resulting solution was stirred at -78 °C for 2 h and the reaction quenched by
- 10 addition of NH₄Cl (sat. sol). After reaching room temperature, the reaction mixture was diluted with Et₂O (50 ml) and washed with brine. The organic phase was dried (MgSO₄) and solvents removed in rotavapor, yielding 7.0 g of crude ester **19**. ¹H NMR (CDCl₃) δ 1.3-2.0 (m, 6H), 3.6-3.9 (m, 2H), 4.45 (s, 2H), 5.27 (s, 2H), 7.40 (m, 5H).
- 15 Step 2: Crude THP-ether **19** (7.0 g) was dissolved in CH₃OH (15 ml) and a catalytic amount of PTSA (250 mg) added at room temperature. After 15 min, the solution was diluted with Et₂O and brine, and the organic phase washed with NaHCO₃ (sat. sol.) and brine, and dried (MgSO₄). Solvents were removed in rotavapor, yielding 4.6 g of crude
- 20 alcohol **20**. ¹H NMR (CDCl₃) δ 4.45 (s, 2H), 5.27 (s, 2H), 7.40 (m, 5H).
- Step 3: Crude alcohol **20** (4.6 g) was dissolved in CH₂Cl₂ (50 ml) containing DMAP (catalytic amounts) and Et₃N (3 ml) at 0 °C. Cinnamoyl chloride (3.3 g, 20 mmol) was added and the mixture stirred at 0 °C for 30 min. The resulting suspension was diluted with Et₂O and
- 25 water, and the organic phase successively washed with NaOH (10%, 50 ml), HCl (2N, 50 ml) and brine, dried (MgSO₄) and the solvents removed in rotavapor, yielding crude ester **21** (7.2 g). ¹H NMR (CDCl₃) δ 5.0 (s, 2H), 5.27 (s, 2H), 6.50 (d, J = 16 Hz, 1H), 7.40-7.70 (m, 10H), 7.80 (d, J = 16 Hz, 1H).

Step 4: A solution of cinnamic ester **21** (7.2 g) in o-xylene (50 ml) was degassed with Ar and heated at 190 °C in pressure tube for 18 h. The mixture was cooled down, and the solvent removed in rotavapor. After chromatographic purification, lactone **22** (3.0g, 44% yield from ether **18**) was obtained. ¹H NMR (CDCl₃) δ 3.62 (m, 1H), 3.78 (d, J = 15.2 Hz, 1H), 4.08 (t, J = 8.8 Hz, 1H), 4.68 (t, J = 8.8 Hz, 1H), 5.34 (d, J = 12.0 Hz, 1H), 5.42 (d, J = 12 Hz, 1H), 7.2-7.5 (m, 10H).

Step 5: Lactone **22** (0.9 g) was dissolved in CH₃OH (40 ml) and hydrogenated under 60 psi of total pressure in the presence of PtO₂ (150 mg) for 14 h. The catalyst was filtered off through a celite bed, and the solvent removed in rotavapor. Acid **23** (270 mg, 41%) crystallizes from EtOAc and hexanes. ¹H-NMR (CDCl₃) δ 2.93 (dd, 3.2; 15.3 Hz, 1H), 3.52 (dd, J = 7.8; 15.3 Hz, 1H), 3.23 (m, 1H), 3.55 (m, 1H), 3.61 (d, J = 3.2 Hz, 1H), 3.7 (dd, J = 5.5; 9.5 Hz, 1H), 4.44 (t, J = 9.2 Hz, 1H), 7.1-7.2 (m, 4H).

Step 6: Carboxylic acid **23** (0.18 g) was suspended in CH₂Cl₂ (5 mL) containing (COCl)₂ (0.15 ml) under an N₂ atmosphere at room temperature. A drop of DMF was added and the resulting mixture stirred at room temperature for 1 h. Solvents were removed in rotavapor, and the crude solid obtained was washed twice with toluene, removing the solvent in rotavapor. A white solid results, which was treated with a solution of toluene (5 ml) and tributyltin hydride (0.3 ml) containing catalytic amounts of Pd(PPh₃)₄ at 0 °C. After 2 h, the mixture was diluted with Et₂O and the organic phase washed with brine.

Chromatographic purification afforded aldehyde **24** (95 mg, 56% yield) as a solid. ¹H-NMR (CDCl₃) δ 2.75 (ddd, 1.0; 7.1; 15.5 Hz, 1H), 3.52 (dd, J = 2.3; 15.5 Hz, 1H), 3.5 (m, 1H), 3.64 (d, J = 1.8 Hz, 1H), 3.78 (m, 1H), 3.83 (dd, J = 4.7; 9.2 Hz, 1H), 4.55 (t, J = 9.1 Hz, 1H), 7.2-7.4 (m, 4H); 9.56 (s, 1H).

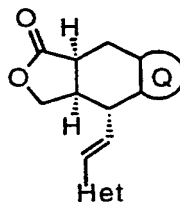
Step 7: A solution of phosphonate **25** (125 mg, 0.45 mmol) in dry THF (10 ml) at 0 °C was treated with *n*-BuLi (0.2 ml, 0.5 mmol, 2.5N in hexanes). After 15 min, a solution of aldehyde **24** (95 mg) in dry THF was added. The resulting solution was stirred at that temperature for 30 min, and diluted with Et₂O and brine. The organic phase was washed with brine and dried (MgSO₄). Solvents are removed in rotavapor, and chromatographic purification affords the title compound (50 mg, 32%).

$^1\text{H-NMR}$ (CDCl_3): δ 8.35 (d, $J = 8.5$ Hz, 1H); 8.0 (d, $J = 9.4$ Hz, 1H); 7.54 (d, $J = 8.6$ Hz, 1H); 7.42 (dd, $J = 2.9; 9.2$, 1H); 7.28-7.36 (m, 4H); 7.11 (d, $J = 2.8$ Hz, 1H); 6.94 (dd, $J = 7.6; 15.9$ Hz, 1H); 6.78 (d, $J = 15.9$ Hz, 1H); 4.56 (dd, $J = 8.3; 9.4$ Hz, 1H); 4.18 (dd, $J = 4.4; 9.4$ Hz, 1H); 4.0 (s, 3H);

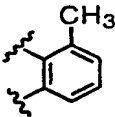
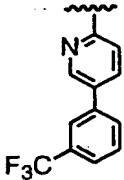
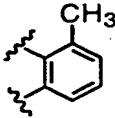
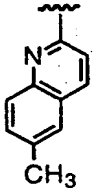
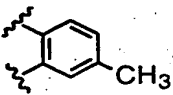
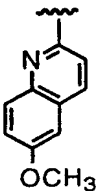
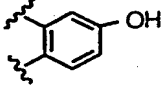
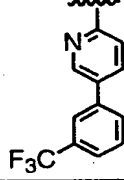
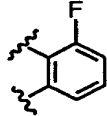
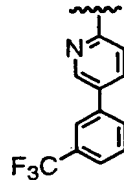
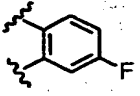
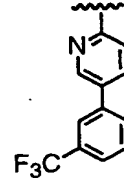
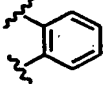
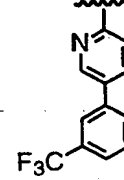
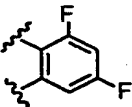
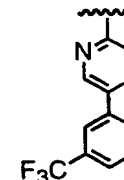
5 3.6 (t, $J = 7.3$ Hz, 1H).

$^{13}\text{C-NMR}$ (CDCl_3): 179.4; 157.6; 152.5; 143.9; 137.3; 135.2; 135.1; 133.9; 133.6; 130.5; 128.3; 127.5; 127.3; 126.9; 122.3; 119.3; 105.0; 71.5; 55.4; 45.6; 39.0; 38.4; 28.7.

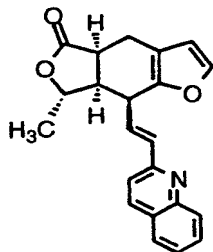
10 Using a similar procedure, compounds of the following structural formulas were prepared, wherein the variables are as defined in the table:



Ex.	Q	Het	Physical Data
3A			MS m/z 430(18), 403(28), 402(100), 401(14)
3B			HRMS (MH^+) found: 386.1756
3C			HRMS (MH^+) found: 466.1625
3D			HRMS (MH^+) found: 402.1709

3E			HRMS (MH+) found: 450.1684
3F			HRMS (MH+) found: 370.1799
3G			HRMS (MH+) found: 386.1750
3H			m.p. 173-176°C
3I			HRMS (MH+) found: 454.1427
3J			HRMS (MH+) found: 454.1423
3K			HRMS (MH+) found: 436.1533
3L			HRMS (MH+) found: 472.1332

Using a similar procedure, the following compound, 3M, is prepared:

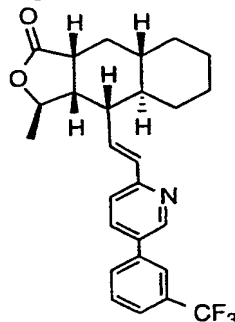


HRMS (MH⁺) found: 346.1445.

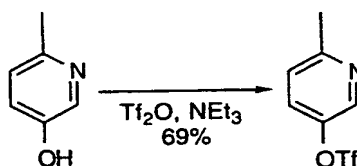
5

EXAMPLE 4

(+)-(3*R*,3*aS*,4*S*,4*aR*,8*aS*,9*aR*)-Decahydro-4-[(*E*)-2-[5-[3-(trifluoromethyl)phenyl]-2-pyridinyl]ethenyl]-3-Methylnaphtho-[2,3-*c*]furan-1(3*H*)-one



Step 1:



10

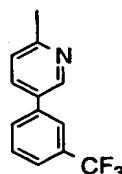
Triflic anhydride (46 ml, 0.275 mol) was added dropwise to a stirred solution of 3-hydroxy-6-methylpyridine (10 g, 0.092 mol) in pyridine (200 ml) at 0 °C and stirred at 0 °C to room temperature for 16 h. The mixture was poured into ice-water (300 ml) and extracted with Et₂O. The Et₂O layer was washed with water (2 x 150 ml) and brine, dried (MgSO₄), and concentrated *in vacuo* to give the desired product (18.7 g, 83%) as a brown oil. ¹H NMR (400 MHz, CDCl₃) δ 2.67 (s, 3H), 7.32 (d, 1H, *J* = 8.5 Hz), 7.57 (dd, 1H, *J* = 8.6, 2.8 Hz), 8.53 (d, 1H, *J* = 2.8 Hz). MS (ESI) *m/z* 242 (MH⁺, 100%); Anal. calc'd for C₇H₆F₃NO₃S: C, 34.86; H, 2.51; N, 5.81. Found: C, 35.24; H, 2.48; N, 5.54.

15

20

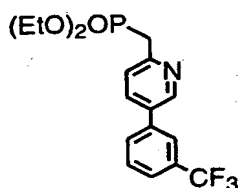
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Step 2:



To a solution of the product of Step 1 (8.5 g, 34.5 mmol) and 3-trifluoromethylphenylboronic acid (10 g, 55 mmol) in toluene (100 ml) was added EtOH (25 ml), K₂CO₃ (14.3 g, 104 mmol) in H₂O (50 ml) and
5 Pd(PPh₃)₄ (400 mg, 0.345 mmol). The mixture was heated in a closed pressure tube under Ar at 120 °C for 16 h. The mixture was diluted with EtOAc, washed with 5% NaOH and brine, dried (MgSO₄), and concentrated *in vacuo*. Flash chromatography of the residue on a silica gel column with EtOAc:hexane (10:90, then 20:80) as eluent gave the
10 desired product (6.7 g, 82%) as yellow solids. ¹H NMR (400 MHz, CDCl₃) δ 2.68 (s, 3H), 7.32 (d, 1H, *J* = 8 Hz), 7.62–7.90 (m, 5H), 8.79 (d, 1H, *J* = 2 Hz). Anal. calc'd for C₁₃H₁₀F₃N•0.10 H₂O: C, 65.32; H, 4.30; N, 5.86. Found: C, 65.27; H, 4.44; N, 5.78.

Step 3:

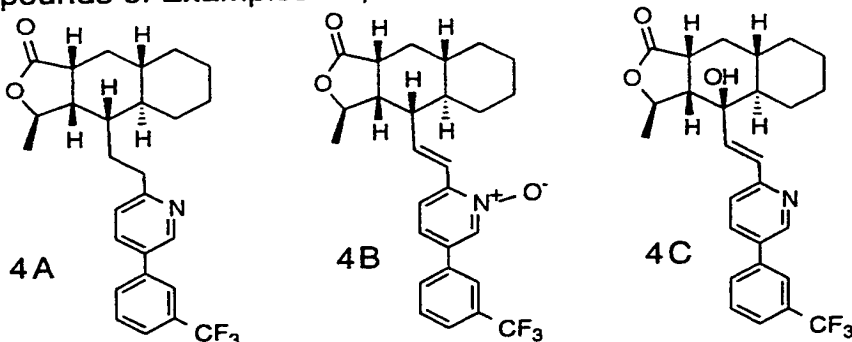


15 Using a procedure similar to that described in Example 1, Step 8, treat the product of Step 2 to obtain the desired product (8.84 g, 85%) as a tan oil. ¹H NMR (400 MHz, CDCl₃) δ 1.36 (t, 6H, *J* = 7 Hz), 3.56 (d, 2H, *J* = 22 Hz), 4.19 (dq, 4H, *J* = 7, 7 Hz), 7.58–7.96 (m, 6H), 8.84 (d, 1H, *J* = 2 Hz); MS (FAB) *m/z* 374 (MH⁺, 100%); Anal. calc'd for
20 C₁₇H₁₉F₃NO₃P•0.25 H₂O: C, 54.04; H, 5.20; N, 3.71. Found: C, 54.22; H, 5.54; N, 3.93.

Step 4: Treat the product of Step 3 in a manner similar to that described in Example 1, Step 9, to obtain the title compound.

25 ¹H NMR (400 MHz, CDCl₃) δ 0.83–2.03 (m, 12H), 1.49 (d, 3H, *J* = 6 Hz), 2.38–2.51 (m, 2H), 2.72–2.81 (m, 1H), 4.79–4.88 (m, 1H), 6.57–6.73 (m, 2H), 7.30–7.95 (m, 6H), 8.85 (d, 1H, *J* = 2 Hz); MS (FAB) *m/z* 456 (MH⁺, 100%). HCl salt: off-white solids; [α]_D²² = +17.0° (c 0.33, MeOH); Anal. calc'd for C₂₇H₂₈F₃NO₂•HCl•0.50 H₂O: C, 64.73; H, 6.04; N, 2.80. Found: C, 64.57; H, 6.32; N, 2.94.

The product of Example 4 is treated as described below to obtain the compounds of Examples 4A, 4B and 4C:

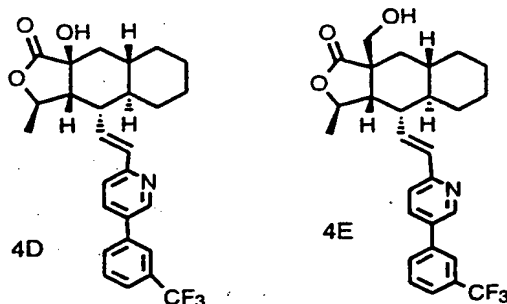


- 5 **4A:** The product of Example 4 (21 mg, 0.050 mmol) was stirred under H₂ (1 atm) over Pd/C (30 mg) in CH₃OH (10 mL) at room temperature for 8 h. The mixture was filtered and the filtrate was concentrated *in vacuo* to obtain 4A (52 mg, 88%) as semi-solids. $[\alpha]^{22}_D = -35.1^\circ$ (*c* 0.69, MeOH); MS (FAB) *m/z* 458 (MH⁺, 100%).
- 10 **4B:** To a solution of the product of Example 4 (20 mg, 0.044 mmol) in CH₂Cl₂ (1 mL) was added m-CPBA (11 mg, 0.066 mmol) at room temperature. The mixture was stirred at room temperature for 16 h. The mixture was diluted with CH₂Cl₂, washed with NaHCO₃ (sat.), dried (MgSO₄), and concentrated *in vacuo*. Preparative TLC separation of the
- 15 residue with CH₂Cl₂:CH₃OH (95:5) as eluent gave 4B (19 mg, 91%) as off-white solids. $[\alpha]^{24}_D = +23.3^\circ$ (*c* 0.43, CH₃OH); MS (ESI) *m/z* 472 (MH⁺, 100%).
- 4C:** The product of Example 4 (21 mg, 0.050 mmol) was heated with SeO₂ (0.2 mg, 0.23 mmol) in 1,4-dioxane (2 mL) at reflux for 40 min.
- 20 The mixture was concentrated *in vacuo*. Preparative TLC separation of the residue with EtOAc:hexane (40:60) as eluent gave 4C (17 mg, 80%) as white solids. $[\alpha]^{20}_D = +42.8^\circ$ (*c* 0.65, CH₃OH); MS (FAB) *m/z* 472 (MH⁺, 100%).

The product of Example 4 is treated as described below to obtain

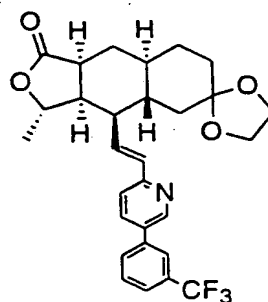
25 the compounds of Example 4D and 4E:

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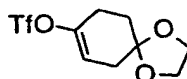
- 4D:** $\text{LiN}(\text{TMS})_2$ (0.6 mL, 0.60 mmol) was added to a solution of the product of Example 4 (227 mg, 0.50 mmol) in dry THF (5 mL) at -78°C and stirred at -78°C for 30 min. and at room temperature for 1 h. A solution of (10-camphorsulfonyl)oxaziridine (137 mg, 0.60 mmol) in dry THF (2 mL) was added at -78°C and stirred at -78°C for 30 min. and at room temperature for 2 h. The mixture was neutralized with saturated NH_4Cl solution and extracted with EtOAc. The organic layer was washed with brine, dried, and concentrated *in vacuo*. Flash chromatography of the residue on a silica gel column with EtOAc:hexane (40:60) as eluent gave 4D (100 mg). MS: 472 (MH^+).
- 4E:** $\text{LiN}(\text{TMS})_2$ (0.6 mL, 0.60 mmol.) was added to a solution of the product of Example 4 (227 mg, 0.50 mmol) in dry THF (5 mL) at -78°C and stirred at -78°C for 30 min. and at room temperature for 1 h. A mixture of paraformaldehyde (225 mg, 2.5 mmol) in dry THF (2 mL) was added at -78°C and stirred at -78°C for 30 min and at room temperature for 2 days. The mixture was neutralized with saturated NH_4Cl solution and extracted with EtOAc. The organic layer was washed with brine, dried, and concentrated *in vacuo*. Flash chromatography of the residue on a silica gel column with EtOAc:hexane (40:60) as eluent gave 4E (30 mg). MS: 486 (MH^+).

EXAMPLE 5



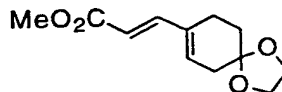
-92-

Step 1:



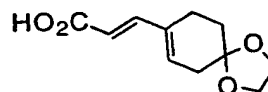
To a solution of 1,4-cyclohexanedione mono-ethylene ketal (10 g, 64 mmol) and 2,6-di-*tert*-butyl-4-methylpyridine (21 g, 102 mmol) in
5 CH_2Cl_2 (350 mL) at room temperature was added triflic anhydride (16 mL, 96 mmol) and the mixture was stirred for 16 h. The mixture was washed with NaHCO_3 (sat.). The organic layer was dried (MgSO_4) and concentrated *in vacuo*. Flash chromatography of the residue on a silica gel column with EtOAc:hexane (5:95 then 10:90) as eluent gave the
10 desired product (13.4 g, 72%) as a clear oil.

Step 2:



To a solution of the product of Step 1 (13 g, 46 mmol) in DMF (150 mL) was added methyl acrylate (8.4 mL, 92 mmol), Et_3N (19 mL, 138 mmol), and $\text{Pd}(\text{PPh}_3)_2\text{Cl}_2$ (1.62 g, 2.3 mmol). The mixture was
15 stirred at 75 °C for 10 h. The mixture was diluted with NH_4Cl (sat.) and extracted with ether. The organic layer was washed with brine, dried (MgSO_4), and concentrated *in vacuo*. Flash chromatography of the residue on a silica gel column with EtOAc:hexane (15:85) as eluent
20 gave the desired product (9.15 g, 89%) as a clear oil.

Step 3:



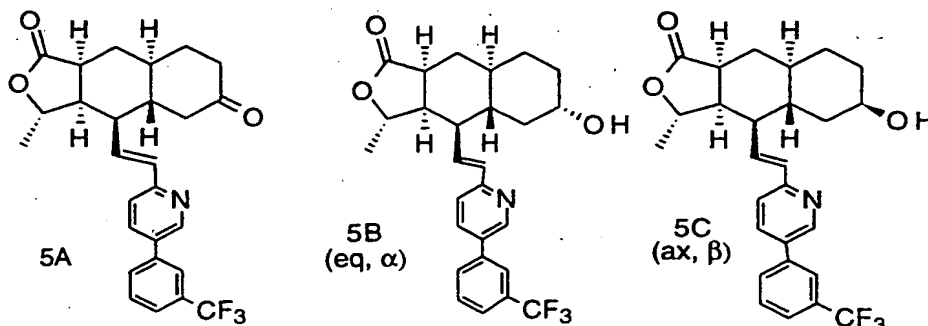
To a solution of the product of Step 2 (9.15 g, 40 mmol) in THF: CH_3OH (450 mL, 1:1) was added NaOH (225 mL, 10%). The mixture
25 was stirred at room temperature for 3 h. The mixture was diluted with water, washed with CH_2Cl_2 , acidified with 10% HCl , and extracted with EtOAc. The organic layer was washed with brine, dried (MgSO_4), and concentrated *in vacuo* to give the desired compound (8.00 g, 95%) as light-yellow solids.

30 Step 4:

The product of Step 3 was treated in a manner similar to that described in Example 1, steps 3 through 6 and 9 to obtain the title

compound (racemate) as off-white solids; MS (ESI) m/z 514 (MH^+ , 100%).

The product of Example 5 is treated as described below to obtain the compounds of Examples 5A, 5B and 5C:



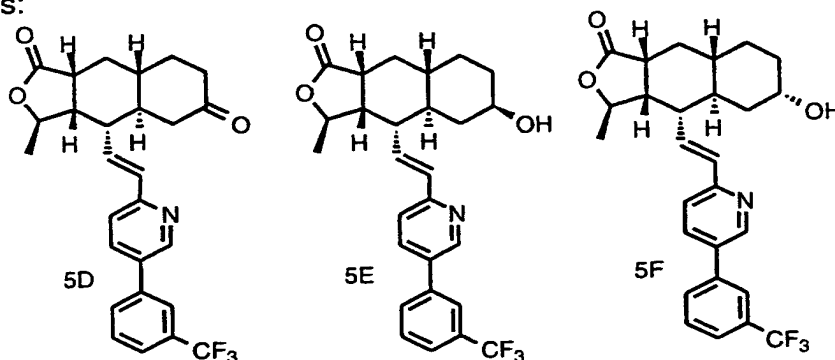
5

5A: A mixture of the compound of Example 5 (65 mg, 0.13 mmol) and HCl (2 mL, 5%) in acetone (2 mL) was stirred at reflux temperature for 16 h. The mixture was neutralized with $NaHCO_3$ (sat.) and extracted with EtOAc. The organic layer was washed with brine, dried ($MgSO_4$) and concentrated *in vacuo*. Preparative TLC separation of the residue with EtOAc:hexane (40:60) as eluent gave 5A (42 mg, 71%), the HCl salt, as white solids. MS (FAB) m/z 470 (MH^+ , 100%).

5B: To a solution of 5A (70 mg, 0.15 mmol) in THF:CH₃OH (10 mL, 1:1) was added $NaBH_4$ (11 mg, 0.30 mmol). The mixture was stirred at room temperature for 10 min. The mixture was diluted with NH_4Cl (sat.) and extracted with EtOAc. The organic layer was washed with brine, dried ($MgSO_4$) and concentrated *in vacuo*. Preparative TLC separation of the residue with EtOAc:hexane (60:40) as eluent gave 5B (39 mg, 55%), the HCl salt, as white solids. MS (FAB) m/z 472 (MH^+ , 100%).

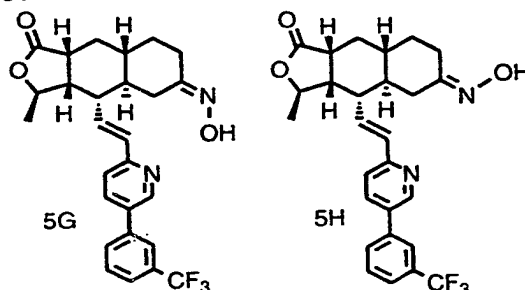
5C: To a solution of 5A (70 mg, 0.15 mmol) in dry THF (5 mL) was added K-Selectride® (0.23 mL, 0.23 mmol, 1.0 M in THF) at -78 °C. The mixture was stirred at -78 °C for 1.5 h. The mixture was diluted with NH_4Cl (sat.) and extracted with EtOAc. The organic layer was washed with brine, dried ($MgSO_4$) and concentrated *in vacuo*. Preparative TLC separation of the residue with EtOAc:hexane (60:40) as eluent gave 5C (45 mg, 63%), the HCl salt, as white solids. MS (FAB) m/z 472 (MH^+ , 100%).

Compounds 5D, 5E and 5F are prepared in a manner similar to that described for Examples 5A–5C using enantiomerically pure starting materials:



5 5D: MS: 470 (MH⁺); 5E: MS: 472 (MH⁺); 5F: MS: 472 (MH⁺)

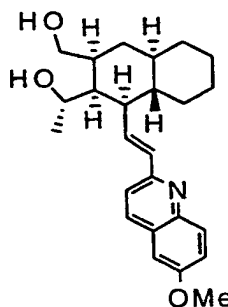
From 5D, the following compounds 5G and 5H can be prepared by conventional methods known to those skilled in the art:



5G: MS: 485 (MH⁺); 5H: MS: 485 (MH⁺).

10

EXAMPLES 6, 6A, 6B, 6C

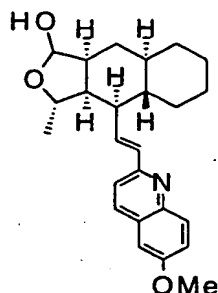


6: To a stirred solution of Example 1AB (0.91 g, 2.3 mmol) in dry THF (20 mL) at room temperature was added LAH (0.18 g, 4.6 mmol) in portions. The mixture was stirred at room temperature for 30 min. NaOH (10%, 0.50 mL) was added dropwise, followed by EtOAc (50 mL) and MgSO₄ powders. The mixture was stirred vigorously for 10 min and

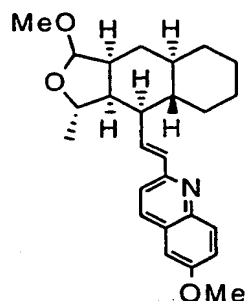
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-95-

filtered. The filtrate was concentrated *in vacuo* to give the title compound (0.88 g, 96%) as white solids. MS (FAB) m/z 396 (MH^+ , 100%).

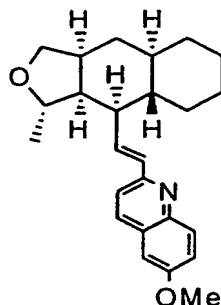


- 5 **6A:** To a solution of Example 1AB (0.200 g, 0.511 mmol) in dry toluene (10 mL) at -78 °C was added DIBAL (0.76 mL, 0.77 mmol, 1.0 M in toluene) dropwise and the mixture was stirred at -78 °C for 1.5 h. EtOAc was added and stirred to room temperature. Water was added and stirred vigorously for 1 h. The organic layer was separated and the aqueous layer was extracted with EtOAc. The combined organic layers were dried ($MgSO_4$) and concentrated *in vacuo*. Flash chromatography of the residue on a silica gel column with EtOAc:hexane (30:70) as eluent gave the title compound (0.100 g, 50%) as white solids. MS (FAB) m/z 394 (MH^+ , 100%).



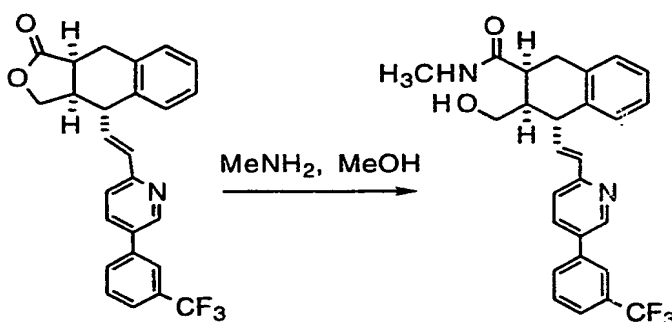
- 15 **6B:** To a solution of Example 6A (50 mg, 0.13 mmol) in CH_3OH (20 mL) at 0 °C was added $BF_3 \cdot OEt_2$ (31 mL, 0.26 mmol) dropwise and the mixture was stirred at 0 °C to room temperature for 3 h. CH_3OH was concentrated *in vacuo*. The residue was partitioned between EtOAc and $NaHCO_3$ (sat.). The organic layer was dried ($MgSO_4$) and concentrated *in vacuo*. Preparative TLC separation of the residue with EtOAc:hexane (20:80) as eluent gave the title compound (30 mg). HCl salt: off-white solids. MS (FAB) m/z 408 (MH^+ , 100%).

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6C: To a solution of Example 6A (50 mg, 0.13 mmol) in CH_2Cl_2 (20 mL) at -78°C was added $\text{BF}_3\cdot\text{OEt}_2$ (31 mL, 0.26 mmol) and Et_3SiH (0.24 mL, 1.3 mmol). The mixture was stirred at -78°C to room temperature for 3 h. The mixture was partitioned between CH_2Cl_2 and NaHCO_3 (sat.). The organic layer was dried (MgSO_4) and concentrated *in vacuo*. Preparative TLC separation of the residue with EtOAc :hexane (20:80) as eluent gave the title compound (38 mg). HCl salt: off-white solids. MS (FAB) m/z 378 (MH^+ , 100%).

EXAMPLES 7, 7A, 7B



7

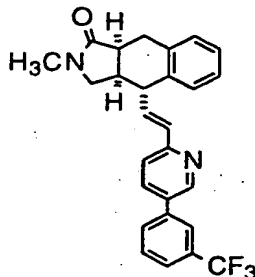
7. The lactone, prepared in a manner similar to Example 3, (75 mg, 0.17 mmol) was dissolved in a 2.0M solution of CH_3NH_2 in CH_3OH (3.0 mL) and stirred under N_2 at room temperature for 1 h. At that time, tlc showed no starting material left, as well as the presence of a more-polar spot. The solution is dissolved in Et_2O (30 mL) and extracted with brine (30 mL x 2), dried (Na_2SO_4) and the solvents removed in rotary evaporator. The title compound (Example 7) was obtained as a white solid (70 mg, 86%). ^1H -NMR (CDCl_3) δ 2.5 (m, 1H); 2.82 (d, $J = 4.6$ Hz, 3H); 3.0 (dd, $J = 5.4$; 16.8 Hz, 1H); 3.07 (m, 1H); 3.25 (dd, $J = 10.1$; 16.8 Hz, 1H); 3.67 (dd, $J = 5.5$; 11.6 Hz, 1H); 3.79 (m, 1H); 3.89 (dd, $J = 8.0$; 11.4, 1H); 6.33 (d, $J = 15.6$ Hz, 1H); 6.49 (m, 1H); 6.95 (dd, $J = 7.0$; 15.6 Hz, 1H); 7.2 (m, 4H); 7.25 (d, $J = 8.2$ Hz, 1H); 7.64 (t, $J = 7.6$ Hz, 1H);

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7.70 (d, J = 7.9, 1H); 7.78 (d, J = 7.6, 1H); 7.82 (m, 2H); 8.73 (d, J = 1.8 Hz, 1H).

¹³C-NMR (CDCl₃): 175.93; 154.80; 147.57; 139.59; 138.22; 134.93; 134.79; 134.24; 133.43; 130.81; 130.57; 130.07; 129.58; 129.11;

5 126.74; 126.30; 124.68; 123.50; 121.74; 62.52; 44.54; 42.89; 39.58; 28.66; 26.40. MS: 467 (M+1). HRMS: 467.1947 (calc.: 467.1946).

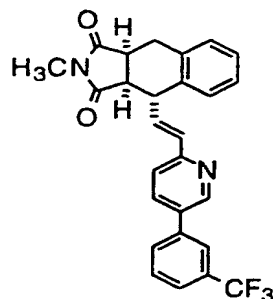


7A: The product of Example 7 (40 mg) was dissolved in dry CH₂Cl₂ (3 mL) under N₂ atmosphere and SOCl₂ (0.1 mL) was added dropwise *via* syringe. The solution was stirred at room temperature for 1 h. The reaction was quenched by addition of NaHCO₃ (sat. sol., 15 mL). The aqueous suspension was extracted with Et₂O (3 x 30 mL). The organic fractions are combined and washed with brine, dried (Na₂SO₄) and taken to dryness in rotary evaporator. The title compound (35 mg, 91%) was obtained as a white solid. ¹H-NMR (CDCl₃): δ 2.82 (m, 1H); 3.0 (dd, J = 6.4; 14.7 Hz, 1H); 3.04 (s, 3H); 3.23 (m, 1H); 3.30 (dd, J = 8.2; 14.7 Hz, 1H); 3.48 (t, 7.9 Hz, 1H); 4.20 (dd, J = 4.4; 9.3 Hz, 1H), 4.48 (dd, J = 7.8; 9.2 Hz, 1H); 6.70 (d, J = 15.6 Hz, 1H); 7.0 (dd, J = 8.4; 15.6 Hz, 1H); 7.25-7.35 (m, 4H); 7.43 (d, J = 7.7 Hz, 1H); 7.64 (t, J = 7.6 Hz, 1H); 7.72 (d, J = 7.6, 1H); 7.83 (d, J = 7.4, 1H); 7.88 (s, broad, 1H); 8.73 (dd, J = 2.4; 8.1 Hz, 1H); 8.84 (d, J = 1.8 Hz, 1H).

¹³C-NMR (CDCl₃): 167.21; 154.36; 148.03; 138.42; 138.18; 136.28; 135.01; 134.89; 133.77; 132.32; 130.15; 129.63; 128.26; 127.07; 126.88; 126.56; 124.74; 124.70; 123.72; 123.68; 121.84; 73.17; 45.12; 40.88; 39.01; 34.18; 30.79.

MS: 449 (M+1). HRMS: 449.1842 (calc.: 449.1841).

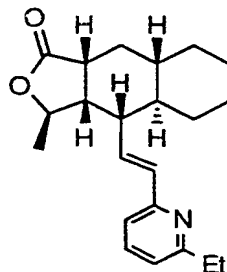
-98-



- 7B: The crude product of Example 7 (45 mg) was dissolved in acetone (2 mL) at 0 °C under N₂ atmosphere. Jones reagent was added dropwise until its red color persisted. The reaction was quenched by
- 5 dropwise addition of ethanol until red color was fully discharged. The resulting green suspension was diluted with Et₂O and water, and the organic phase washed with brine, dried (MgSO₄) and the solvents removed in rotavapor. The crude acid was dissolved in dry CH₂Cl₂ (5 mL) under N₂ atmosphere and SOCl₂ (0.1 mL) added dropwise *via*
- 10 syringe. The solution was stirred at room temperature for 1 h. The reaction was quenched by addition of NaHCO₃ (sat. sol., 15 mL). The aqueous suspension was extracted with Et₂O (3 x 30 mL). The organic fractions were combined and washed with brine, dried (Na₂SO₄) and taken to dryness in rotary evaporator. After flash chromatography (30%
- 15 EtOAc in hexanes) the desired imide (35 mg, 77%) was obtained.
- ¹H-NMR (CDCl₃): δ 2.86 (s, 3H); 3.20 (m, 2H); 3.5 (m, 1H); 3.58 (dd, J = 1.8; 9.2 Hz, 1H); 4.4 (m, 1H); 6.38 (dd, J = 1.2; 15.6 Hz, 1H); 7.10 (dd, J = 5.2; 15.6 Hz, 1H); 7.2-7.4 (m, 5H); 7.65 (t, J = 7.7 Hz, 1H); 7.72 (d, J = 7.6, 1H); 7.8 (d, J = 7.6, 1H); 7.86 (s, 1H); 7.92 (d, J = 8.2 Hz, 1H); 8.86
- 20 (s, 1H). MS: 463 (M+1). HRMS: 463.1634 (calc.: 463.1633).

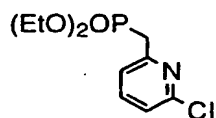
EXAMPLE 8

(+)-(3*R*,3*aS*,4*S*,4*aR*,8*aS*,9*aR*)-Decahydro-4-[(*E*)-2-(6-ethyl-2-pyridinyl)ethenyl]-3-methylnaphtho[2,3-*c*]furan-1(3*H*)-one



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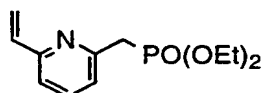
Step 1: Preparation of



Using a procedure similar to that of Example 1, Step 8, 2-chloro-6-methylpyridine was treated with diethylchlorophosphate to obtain a clear oil.

¹H NMR (400 MHz, CDCl₃) δ 1.34 (t, 6H, *J* = 7 Hz), 3.43 (d, 2H, *J* = 22 Hz), 4.15 (dq, 4H, *J* = 7, 7 Hz), 7.27 (dd, 1H, *J* = 8, 2 Hz), 7.38 (dd, 1H, *J* = 8, 2 Hz), 7.66 (t, 1H, *J* = 8 Hz); MS (FAB) *m/z* 264 (MH⁺, 100%).

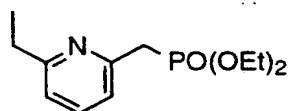
Step 2: Preparation of



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To a solution of the product of Step 1 (5.24 g, 19.9 mmol) in anhydrous THF (100 mL) was added Pd(PPh₃)₄ (1.2 g, 1.0 mmol) and vinyltributyltin (8.72 mL, 29.9 mmol). The mixture was heated in a closed pressure tube under Ar at 120 °C for 16 h. The aqueous layer was neutralized with 10% NaOH and solid NaHCO₃ and extracted with CH₂Cl₂. The organic layer was dried (MgSO₄) and concentrated *in vacuo*. Flash chromatography of the residue on a silica gel column with EtOAc:hexane (40:60 then 80:20) as eluent gave the desired product (3.66 g, 72%) as a clear oil. ¹H NMR (400 MHz, CDCl₃) δ 1.32 (t, 6H, *J* = 7 Hz), 3.48 (d, 2H, *J* = 22 Hz), 4.15 (dq, 4H, *J* = 7, 7 Hz), 5.52 (d, 1H, *J* = 11 Hz), 6.26 (d, 1H, *J* = 17 Hz), 6.85 (dd, 1H, *J* = 17, 11 Hz), 7.26–7.34 (m, 2H), 7.66 (t, 1H, *J* = 8 Hz); MS (CI) *m/z* 256 (MH⁺, 100%); Anal. calc'd for C₁₂H₁₈NO₃P•0.50 H₂O: C, 54.54; H, 7.25; N, 5.30; P, 11.72. Found: C, 54.80; H, 7.21; N, 5.34; P, 11.87.

25 Step 3: Preparation of



To a stirred solution of the product of Step 2 (3.58 g, 14.0 mmol) in CH₃OH (100 mL) was added 5% Pd/C (0.36 g). The mixture was stirred under H₂ (1 atm) at room temperature for 16 h. The mixture was diluted with EtOAc and extracted with 10% HCl. The solids were filtered and washed with CH₃OH. The filtrate and washings were combined

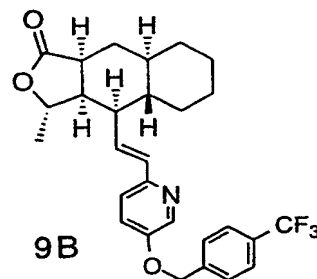
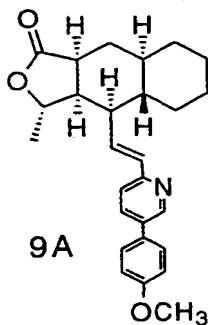
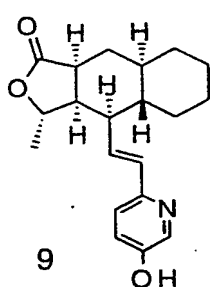
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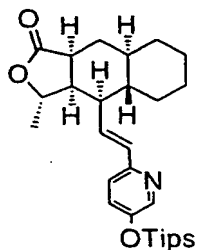
and concentrated *in vacuo* to give the desired product (3.56 g, 99%) as a clear oil. ^1H NMR (400 MHz, CDCl_3) δ 1.32 (t, 6H, $J = 7$ Hz), 1.34 (t, 3H, $J = 7.6$ Hz), 2.84 (q, 2H, $J = 7.6$ Hz), 3.44 (d, 2H, $J = 22$ Hz), 4.13 (dq, 4H, $J = 7, 7$ Hz), 7.08 (d, 1H, $J = 7.6$ Hz), 7.25 (d, 1H, $J = 7.6$ Hz), 7.59 (t, 1H, $J = 7.6$ Hz); MS (CI) m/z 258 (MH^+ , 100%); Anal. calc'd for $\text{C}_{12}\text{H}_{20}\text{NO}_3\text{P} \cdot 0.50 \text{H}_2\text{O}$: C, 54.13; H, 7.95; N, 5.26; P, 11.63. Found: C, 54.19; H, 7.95; N, 5.25; P, 11.65.

Step 4: Using a procedure similar to that described in Example 1, Step 9, combine the product of Step 3 with the product of Example 1, Step 6, to obtain the title compound as a white gum. ^1H NMR (400 MHz, CDCl_3) δ 0.78–2.01 (m, 12H), 1.36 (t, 3H, $J = 7.6$ Hz), 1.49 (d, 3H, $J = 6$ Hz), 2.36–2.43 (m, 2H), 2.70–2.78 (m, 1H), 2.86 (q, 2H, $J = 7.6$ Hz), 4.77–4.85 (m, 1H), 6.47–6.58 (m, 2H), 7.06 (d, 1H, $J = 7.6$ Hz), 7.11 (d, 1H, $J = 7.6$ Hz), 7.59 (t, 1H, $J = 7.6$ Hz). HCl salt: off-white solids; $[\alpha]^{22}_{\text{D}} = +21.3^\circ$ (c 0.41, CH_3OH); MS (ESI) m/z 340 (MH^+ , 100%). Anal. calc'd for $\text{C}_{22}\text{H}_{29}\text{NO}_2 \cdot \text{HCl} \cdot 1.50 \text{H}_2\text{O}$: C, 65.58; H, 8.25; N, 3.48. Found: C, 65.54; H, 8.40; N, 3.68.

EXAMPLES 9, 9A and 9B



- 20 **Example 9, Step 1:** Using a procedure similar to that of Example 1, Step 7, treat 3-hydroxy-6-methylpyridine with triisopropylsilyl chloride.
- Step 2:** Using a procedure similar to that of Example 1, Step 8, treat the product of Step 1 with diethylchlorophosphate.
- Step 3:**

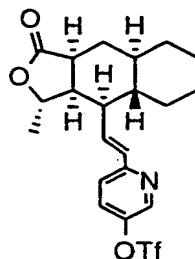


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Using a procedure similar to that of Example 1, Step 9, combine the product of Step 3 with the product of Example 1, Step 6, to obtain the desired product (Tips is triisopropylsilyl) as white solids; MS (FAB) m/z 484 (MH^+ , 100%).

- 5 Step 4: By treating the product of Step 3 as described in Example 1, Step 10, the product of Example 9 was obtained. HCl salt, off-white solids; MS (CI) m/z 342 (MH^+ , 100%).

Example 9A, Step 1:



- 10 To a solution of the product of Example 9 (30 mg, 0.092 mmol) and Et_3N (64 mL, 0.46 mmol) in CH_2Cl_2 (5 mL) at room temperature was added triflic anhydride (46 mL, 0.28 mmol) and the mixture was stirred for 10 min, then washed with water. The organic layer was dried ($MgSO_4$) and concentrated *in vacuo*. Flash chromatography of the residue on a silica gel column with $EtOAc:hexane$ (40:60) as eluent
15 gave the desired triflate (42 mg, 100%). HCl salt, light-yellow solids. MS (FAB) m/z 460 (MH^+ , 100%).

- Step 2: To a solution of the product of Step 1 (37 mg, 0.081 mmol) and *p*-methoxyphenylboronic acid (24 mg, 0.16 mmol) in toluene (2 mL) was
20 added $EtOH$ (0.5 mL), K_2CO_3 (44 mg, 0.32 mmol) in H_2O (1 mL), and $Pd(PPh_3)_4$ (9 mg, 0.008 mmol). The mixture was heated in a closed pressure tube under Ar at 120 °C for 16 h. The mixture was diluted with $EtOAc$, washed with 5% NaOH and brine, dried ($MgSO_4$) and concentrated *in vacuo*. Preparative TLC separation of the residue with
25 $EtOAc:hexane$ (40:60) as eluent gave 9A (24 mg, 71%). HCl salt, white solids. MS (CI) m/z 418 (MH^+ , 100%).

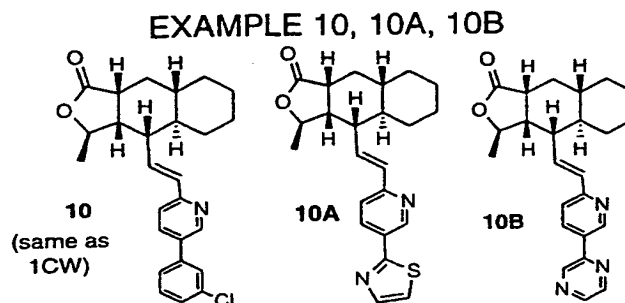
Example 9B:

- A mixture of the product of Example 9 (33 mg, 0.10 mmol), 4-(trifluoromethyl)benzyl bromide (36 mg, 0.15 mmol), and K_2CO_3 (42 mg,
30 0.30 mmol) in acetone (2 mL) was stirred at reflux temperature for 3 h. The solids were filtered and washed with $EtOAc$. The filtrate and the

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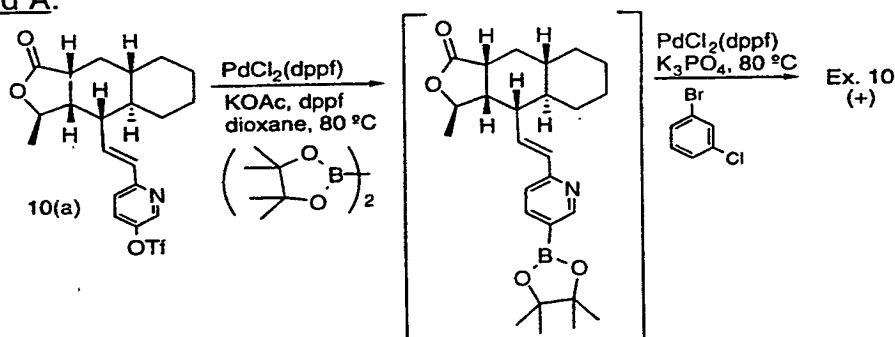
washings were combined and concentrated *in vacuo*. Preparative TLC separation of the residue with EtOAc:hexane (40:60) as eluent gave Example 9B (41 mg, 85%). HCl salt, off-white solids. MS (CI) m/z 486 (MH⁺, 100%).

5



Alternatives to coupling procedure of Example 9:

Method A:



- 10 A mixture of 10(a) (prepared similarly to Example 9A, step 1, using appropriate starting material, 460 mg, 1.00 mmol), diboron pinacol ester (305 mg, 1.20 mmol), potassium acetate (294 mg, 3.00 mmol), 1,1'-bis(diphenylphosphino)ferrocene (55 mg, 0.10 mmol), and dichloro[1,1'-bis(diphenylphosphino)ferrocene]palladium (II) dichloromethane adduct (82 mg, 0.10 mmol) in 1,4-dioxane (5 mL) was heated in a closed pressure tube under N₂ at 80 °C for 2 h. The mixture was cooled to room temperature. To this mixture was added 1-bromo-3-chlorobenzene (235 μ L, 2.00 mmol), K₃PO₄ (636 mg, 3.00 mmol), dichloro-[1,1'-bis(diphenylphosphino)ferrocene]-palladium (II) dichloromethane adduct (41 mg, 0.050 mmol), and 1,4-dioxane (5 mL). The mixture was heated in a closed pressure tube under nitrogen at 80 °C for 16 h. The mixture was partitioned between NH₄Cl (saturated) and EtOAc. The organic layer was washed with brine, dried (MgSO₄), and concentrated *in vacuo*. Flash chromatography of the residue on a silica gel column with
- 15
- 20

-103-

EtOAc:hexane (20:80, then 25:75) as eluent gave (+)-Example 10 (360 mg, 85%) as off-white solids. HCl salt: off-white solids; MS (FAB) m/z 422 (MH^+ , 100).

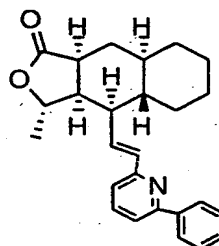
Method B:

- 5 10A: A mixture of 10(a) (46 mg, 0.10 mmol), 2-tributylstannylthiazole (112 mg, 0.30 mmol), and tetrakis(triphenylphosphine)palladium (12 mg, 0.010 mmol) in N-methylpyrrolidinone (1 mL) was heated in a closed pressure tube under nitrogen at 120 °C for 20 h. The mixture was partitioned between H₂O and ether. The organic layer was washed with
10 brine, dried (MgSO₄), and concentrated *in vacuo*. Preparative TLC of the residue on a silica gel plate with EtOAc:hexane (30:70) as eluent gave Example 10A (17 mg). MS: 395 (MH^+).

10B: Example 10B was prepared using a similar procedure:
MS: 392 (MH^+).

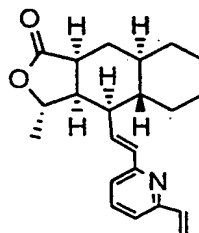
15

EXAMPLE 11



- To a solution of Example 1L (20 mg, 0.058 mmol) and phenylboronic acid (14 mg, 0.12 mmol) in toluene (2 mL) was added EtOH (0.5 mL), K₂CO₃ (32 mg, 0.23 mmol) in H₂O (1 mL), and
20 Pd(PPh₃)₄ (7 mg, 0.006 mmol). The mixture was heated in a closed pressure tube under Ar at 120 °C for 16 h. The mixture was diluted with EtOAc, washed with 5% NaOH and brine, dried (MgSO₄), and concentrated *in vacuo*. Preparative TLC separation of the residue with EtOAc:hexane (20:80) as eluent gave the title compound (10 mg, 71%).
25 HCl salt, off-white solids. MS (CI) m/z 388 (MH^+ , 100%).

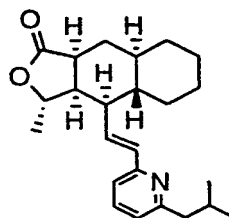
EXAMPLE 12



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Example 1L (333 mg, 0.963 mmol) was heated with vinyltri-*n*-butyltin (424 μ L, 1.44 mmol) and Pd(PPh₃)₄ (62 mg, 0.05 mmol) in THF (10 mL) in a closed pressure tube under Ar at 120 °C for 16 h. The mixture was diluted with EtOAc, washed with NH₄Cl (sat.) and brine, dried (MgSO₄) and concentrated *in vacuo*. Flash chromatography of the residue on a silica gel column with EtOAc:hexane (20:80) as eluent gave the title compound (281 mg, 86%) as white solids. MS (CI) *m/z* 338 (MH⁺, 100%).

EXAMPLE 13

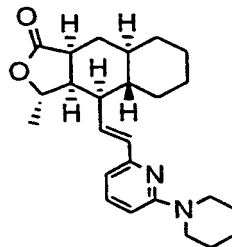


10

To a solution of ZnCl₂ (0.96 mL, 0.5 M in THF) was added isobutylmagnesium chloride (0.22 mL, 2.0 M in ether) at -78 °C. The mixture was stirred at -78 °C to room temperature for 1 h. Example 1L (30 mg) and Pd(PPh₃)₄ (10 mg) were added to the resulting mixture. The mixture was heated in a closed pressure tube under Ar at 120 °C for 2.5 h. The mixture was diluted with EtOAc, washed with NH₄Cl (sat.), dried (MgSO₄) and concentrated *in vacuo*. Preparative TLC separation of the residue with EtOAc:hexane (20:80) as eluent gave the title compound (16 mg) as the HCl salt, white solids. MS (FAB) *m/z* 368 (MH⁺, 100%).

20

EXAMPLE 14



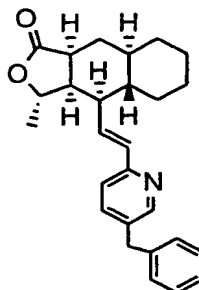
Example 1L (20 mg, 0.058 mmol) was heated with piperidine (0.5 mL) in a closed pressure tube under Ar at 190 °C for 13 h. The mixture was diluted with EtOAc, washed with NaHCO₃ (sat.) and brine, dried (MgSO₄), and concentrated *in vacuo*. Preparative TLC separation of the

25

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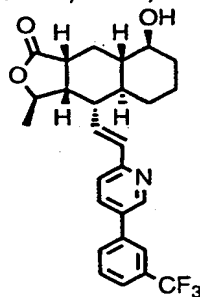
residue with EtOAc:hexane (30:70) as eluent gave the title compound (15 mg, 66%). HCl salt, white solids. MS (CI) m/z 395 (MH^+ , 100%).

EXAMPLE 15

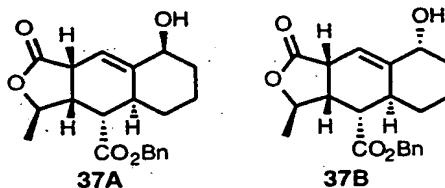


- 5 To a solution of $ZnCl_2$ (0.95 mL, 0.44 mmol, 0.5 M in THF) was added benzylmagnesium chloride (0.44 mL, 0.44 mmol, 1.0 M in ether) at $-78^\circ C$. The mixture was stirred at $-78^\circ C$ to room temperature for 1 h. The product of Example 9A, Step 1 (40 mg, 0.087 mmol) and $Pd(PPh_3)_4$ (10 mg, 0.009 mmol) were added to the resulting mixture. The mixture
10 was heated in a closed pressure tube under Ar at $120^\circ C$ for 16 h. The mixture was diluted with EtOAc, washed with brine, dried ($MgSO_4$) and concentrated *in vacuo*. Preparative TLC separation of the residue with EtOAc:hexane (30:70) as eluent gave the title compound (34 mg, 97%). HCl salt, off-white solids. MS (FAB) m/z 402 (MH^+ , 100%).

15 EXAMPLES 16, 16A, 16B and 16C



Step 1:

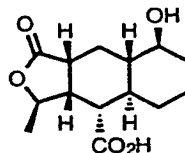


- 20 A mixture of 6 (3.15 g) and SeO_2 (3.10 g) in 1,4-dioxane (50 mL) and pyridine (5 mL) was heated in a closed pressure tube at $100^\circ C$ for 1 h. The mixture was cooled to room temperature, filtered, and

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concentrated *in vacuo*. Flash chromatography of the residue on a silica gel column with EtOAc:hexane (30:70) as eluent gave **37B** (950 mg) and **37A** (1.05 g).

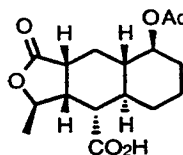
Step 2:



5

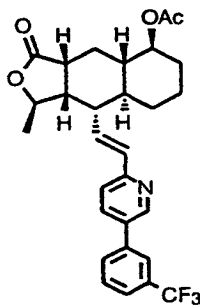
A mixture of **37A** (1.05 g) and PtO_2 (250 mg) in EtOAc (70 mL) was stirred under a hydrogen balloon at room temperature for 16 h. The mixture was filtered and the filtrate was concentrated *in vacuo* to give the desired product (670 mg, 85%).

10 Step 3:



15 A mixture of the product of Step 2 (670 mg) and Ac_2O (2 mL) in pyridine (5 mL) was stirred at room temperature for 16 h. The mixture was poured into a mixture of dilute HCl solution and ice and stirred for 1 h. The resulting mixture was extracted with ether. The organic layer was washed with brine, dried, and concentrated *in vacuo* to give the desired product (700 mg).

Step 4:



20 Treat the product of Step 3 in a manner similar to that described in Example 1, Steps 6 and 9, using the appropriate phosphonate to obtain the desired compound.

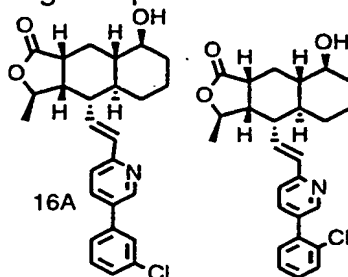
Step 5:

25 A mixture of the product of Step 4 (100 mg), NaOH (10%, 2 mL), and CH_3OH (2 mL) in THF (7 mL) was stirred at 0 °C for 3 h. The

-107-

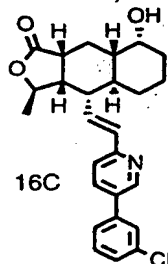
- mixture was neutralized with saturated NH_4Cl solution and extracted with ether. The organic layer was washed with brine, dried, and concentrated *in vacuo*. Flash chromatography of the residue on a silica gel column with EtOAc:hexane (45:55, then 50:50) as eluent gave the
- 5 title product (25 mg). MS (FAB) m/z 472 (MH^+ , 100).

Using the appropriate phosphonates in the procedure of Example 16, Steps 4–5, the following compounds 16A and 16B were prepared:



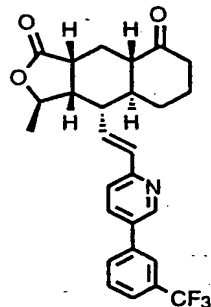
- 10 16A: (MS (ESI) m/z 438 (MH^+ , 100);
- 16B: MS (ESI) m/z 438 (MH^+ , 100).

Using the procedure of Example 16, Steps 2–5, employing starting material **37B**, the following compound 16C was prepared:



MS (FAB) m/z 472 (MH^+ , 100).

EXAMPLES 17 and 17A



15

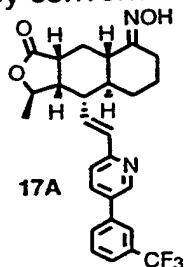
Jones reagent was added to a solution of the product of Example 16 hydrochloride (20 mg) in acetone (5 mL) at room temperature until red color persisted. The reaction was quenched with EtOH and extracted with ether. The organic layer was washed with brine, dried,

-108-

and concentrated *in vacuo* to give the title compound (20 mg).

¹H MS (FAB) *m/z* 470 (MH⁺, 100).

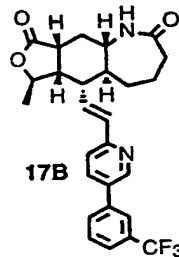
From the compound of Example 17, the following compound can be prepared by conventional methods known to those skilled in the art:



MS (ESI) *m/z* 485 (MH⁺, 100).

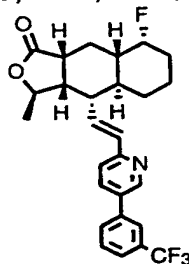
5

From 17A, the following compound can be prepared by conventional methods known to those skilled in the art:



MS (FAB) *m/z* 485 (MH⁺, 100).

EXAMPLES 18, 18A, 18B, 18C, and 18D



10

DAST (diethylaminosulfurtrifluoride) (2–3 drops) was added to a solution of the product of Example 16 (12 mg) in CH₂Cl₂ (2 mL) at 0 °C and the mixture was allowed to warm to room temperature. The mixture was washed with NaHCO₃ solution (sat'd) and brine, dried and concentrated *in vacuo*. Flash chromatography of the residue on a silica gel column with EtOAc:hexane (35:65) as eluent gave the title product (8 mg). MS (ESI) *m/z* 474 (MH⁺, 100).

15

Using the procedure of Example 18, employing as starting material the product of Example 16C, the compound 18A was prepared; similarly, using the appropriate starting materials, compounds 18B, 18C and 18D were also prepared:

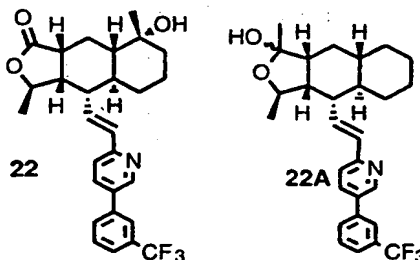
20

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(30:70) as eluent gave the title product (13 mg).

MS (FAB) m/z 490 (MH^+ , 100).

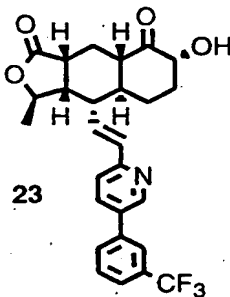
EXAMPLE 22, 22A



- 5 22: MeMgBr (0.1 mL, 1.4 M) was added to a solution of the product of Example 17 (50 mg) in dry THF (3 mL) and stirred at room temperature for a few minutes. The mixture was neutralized with saturated NH_4Cl solution and extracted with ether. The organic layer was washed with brine, dried, and concentrated *in vacuo*. Flash chromatography of the residue on a silica gel column with EtOAc:hexane (40:60) as eluent
10 gave the title product (10 mg). MS (FAB) m/z 486 (MH^+ , 100).

22A: Using the product of Example 4 in a similar procedure, 22A was prepared. MS (FAB) m/z 454 (MH^+ , 100).

EXAMPLES 23, 23A, 23B



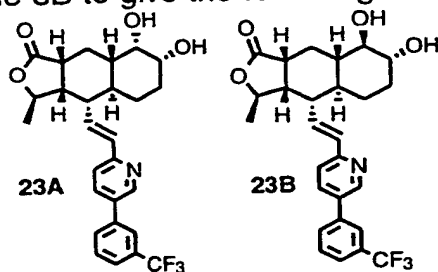
15

- n-BuLi (0.15 mL, 0.22 mmol) was added to a solution of diisopropylamine (0.060 mL, 0.22 mmol) in dry THF at $-78^\circ C$ and stirred for a few minutes. A solution of the product of Example 17 (40 mg, 0.10 mmol) in dry THF (2 mL) was added at $-78^\circ C$ and stirred for 15 min. A
20 solution of (10-camphorsulfonyl)oxaziridine (46 mg, 0.20 mmol) in dry THF (2 mL) was added at $-78^\circ C$ and stirred to room temperature (2 h). The mixture was neutralized with saturated NH_4Cl solution and extracted with ether. The organic layer was washed with brine, dried, and concentrated *in vacuo*. Flash chromatography of the residue on a

-112-

silica gel column with EtOAc:hexane (60:40, then 70:30) as eluent gave the title product (10 mg). MS (ESI) m/z 486 (MH^+ , 100).

The product of Example 23 is treated in a manner similar to that described for Example 5B to give the following compounds:

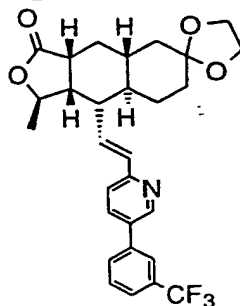


5

23A: MS (FAB) m/z 488 (MH^+ , 100).

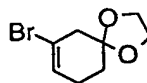
23B: MS (FAB) m/z 488 (MH^+ , 100).

EXAMPLE 24



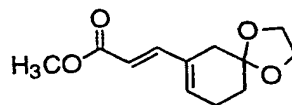
10

Step 1:



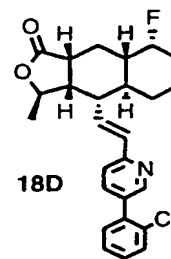
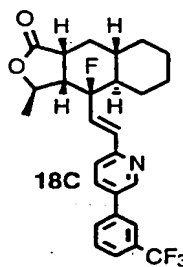
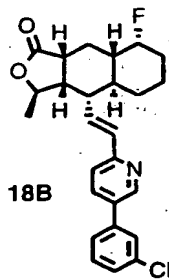
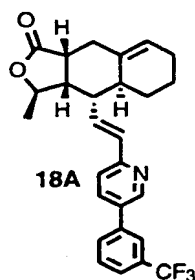
See J. Organometallic Chem., 521 (1996) pp203-210; J. Org. Chem., 47 (1982), pp2825-2832.

15 Step 2:



20

The product of Step 1 (27.5 g, 0.1255 mol) was dissolved in DMF (400 mL) and methylacrylate (23 mL, 0.251 mol), Et_3N (52.25 mL, 0.3765 mol) and $Pd(Ph_3P)_2Cl_2$ (4.37 g, 5 mol %) were added successively. The mixture was heated at $75^\circ C$ for 16 h. The reaction mixture was combined with NH_4Cl (sat'd), extracted with ether and dried ($MgSO_4$). The extracts were concentrated *in vacuo* and the residue



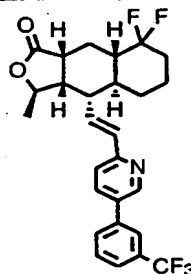
18A: MS (FAB) m/z 454 (MH^+ , 100).

18B: MS (ESI) m/z 440 (MH^+ , 100).

18C: MS (ESI) m/z 474 (MH^+ , 100).

5 18D: MS (ESI) m/z 440 (MH^+ , 100).

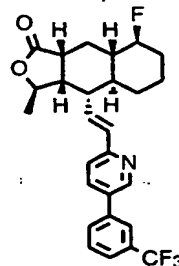
EXAMPLE 19



10 DAST (250 mL) was added to a solution of the product of Example 17 (60 mg) in CH_2Cl_2 (5 mL) and stirred at room temperature for 3 days. The mixture was washed with saturated $NaHCO_3$ solution and brine, dried, and concentrated *in vacuo*. Flash chromatography of the residue on a silica gel column with EtOAc:hexane (35:65) as eluent gave the title product. MS (FAB) m/z 492 (MH^+ , 100).

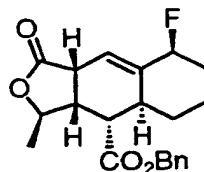
15

EXAMPLES 20, 20A and 20B



Step 1:

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DAST was added to a solution of compound **37B** (180 mg) in CH_2Cl_2 at -78°C and stirred for 15 min. The mixture was washed with saturated NaHCO_3 solution and brine, dried, and concentrated *in vacuo*.

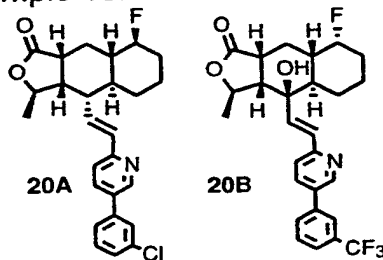
- 5 Flash chromatography of the residue on a silica gel column with EtOAc:hexane (10:90) as eluent gave the desired product (100 mg).

Step 2:

Treat the product of Step 1 in a manner similar to that described in Example 16, Steps 2 and 4, gave the title compound.

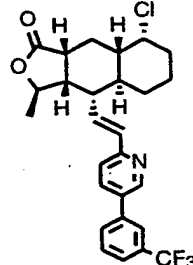
- 10 MS (ESI) m/z 474 (MH^+ , 100).

Using the appropriate phosphonate in the above procedure, **20A** is prepared; using the procedure of **4C**, the compound **20B** is prepared from the product of Example 18:



- 15 20A: MS (CI) m/z 440 (MH^+ , 100).
20B: MS (ESI) m/z 490 (MH^+ , 100).

EXAMPLE 21

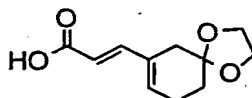


- 20 The product of Example 16 (30 mg, 0.063 mmol) was refluxed in SOCl_2 (1 mL) for 3 h. The mixture was concentrated *in vacuo*. Preparative TLC of the residue on a silica gel plate with EtOAc:hexane

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chromatographed (9:1-4:1 hexane/EtOAc) to give 20 g (71 %) of the desired compound. ^1H NMR (CDCl_3) δ 1.78 (t, $J = 6.5$ Hz, 2H), 2.38 (s, 2H), 2.44 (m, 2H), 3.74 (s, 3H), 4.0 (s, 4H), 5.73 (d, $J = 15$ Hz, 1H), 6.17 (br s, 1H), 7.36 (d, $J = 15$ Hz, 1H).

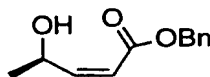
5 Step 3:



The product of Step 2 (20 g, 0.089 mol) was dissolved in a 1:1 mixture of THF/ CH_3OH (520 mL total). 1M NaOH solution (260 mL) was added slowly. The mixture was stirred for 4 h and water was added.

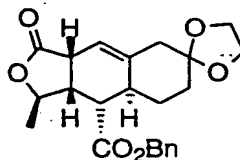
10 The mixture was washed with ether, the aqueous layer was acidified to pH 1 and extracted with EtOAc (x3), the combined extracts were dried (MgSO_4) and the solution concentrated *in vacuo* to give 19 g (99 %) of the desired compound. ^1H NMR (CDCl_3) δ 1.79 (t, $J = 6.5$ Hz, 2H), 2.40 (s, 2H), 2.46 (m, 2H), 4.01 (m, 4H), 5.73 (d, $J = 15.7$ Hz, 1H), 6.23 (s, 1H),
15 7.41 (d, $J = 15.7$ Hz, 1H).

Step 4:



The product of Example 1, Step 2 (23.28 g, 0.114 mol) was dissolved in THF (232 mL) and Lindlar's hydrogenation catalyst was
20 added (3.48 g). The mixture was placed under 1 atm pressure of H_2 (g) and stirred for 2.5 h. The mixture was filtered and concentrated *in vacuo* to give the desired compound (22 g, 93 %). ^1H NMR (CDCl_3) δ 1.32 (d, $J = 6.5$ Hz, 3H), 5.09 (m, 1H), 5.17 (s, 2H), 5.86 (d, $J = 11.7$ Hz, 1H), 6.30 (dd, $J = 11.7, 7.0$ Hz, 1H), 7.38 (s, 5H).

25 Step 5:

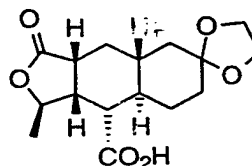


The product of step 3 (18 g, 0.0856 mol) was dissolved in CH_2Cl_2 (350 mL) and cooled to 0°C . 1,3-Dicyclohexylcarbodiimide (23.23 g, 0.112 mol) was added, followed by 4-pyrrolidinopyridine (1.39 g, 9.4
30 mmol). After stirring 5 min, a solution of the product of step 4 (22 g, 0.1067 mol) in CH_2Cl_2 (127 mL) was added over 10 min. The mixture

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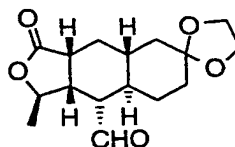
was stirred at 0°C for 2 h and at room temperature for 1 h. The mixture was then filtered, concentrated *in vacuo*, and the residue column chromatographed (9:1-4:1 hexane/EtOAc as eluent) to obtain 27 g of an oil. This product was dissolved in xylene (300 mL) and heated at 215°C for 7 h. Column chromatography (9:1-4:1-2:1 hexane/EtOAc) gave 13.2 g of an oil. The oil was dissolved in THF (264 mL) and DBU (4.9 mL, 0.033 mol) was added. The mixture was stirred for 1 h, diluted with EtOAc (500 mL), washed with NH₄Cl(sat), dried (MgSO₄), concentrated *in vacuo*, filtered through a pad (1 inch) of SiO₂ (eluting with EtOAc) and concentrated *in vacuo* to give the desired compound (13 g, 38 %). ¹H NMR (CDCl₃) δ 1.10 (d, *J* = 6.0 Hz, 3H), 1.2 (m, 1H), 1.65-1.85 (m, 2H), 1.92 (m, 1H), 2.35 (m, 1H), 2.47 (m, 1H), 2.59 (dd, *J* = 10.75, 4.0 Hz, 1H), 2.70 (m, 1H), (q, *J* = 2.5 Hz, 1H), 3.85-4.0 (m, 5H), 4.45 (m, 1H), 5.15 (AB quartet, *J* = 12.0, 10.5 Hz, 2H), 5.36 (br s, 1H), 7.35 (s, 5H).

Step 6:



The product of step 5 (4.92 g, 0.0123 mol) was dissolved in EtOAc (250 mL), 10% palladium on carbon (492 mg) was added and the mixture stirred under 1 atm of H₂ (g) for 1 h. The mixture was filtered through a pad of celite, PtO₂ (492 mg) was added to the filtrate and the mixture stirred for 16 h under 1 atm H₂ (g). The mixture was then filtered and concentrated *in vacuo* to give 3.81 g (99 %) of the desired compound. ¹H NMR (CDCl₃) δ 1.25 (m, 2H), 1.35 (d, *J* = 6.5 Hz, 3H), 1.3-1.5 (m, 3H), 1.6 (m, 1H), 1.7-1.95 (m, 3H), 2.5 (m, 1H), 2.58 (m, 1H), 2.68 (m, 1H), 3.95 (m, 5H), 4.69 (m, 1H).

Step 7:



The product of step 6 (1 g, 3.2 mmol) was dissolved in toluene (20 mL), SOCl₂ (1.25 mL) was added and the mixture heated at 80°C for 16 h. The mixture was concentrated *in vacuo*, dissolved in fresh toluene (16

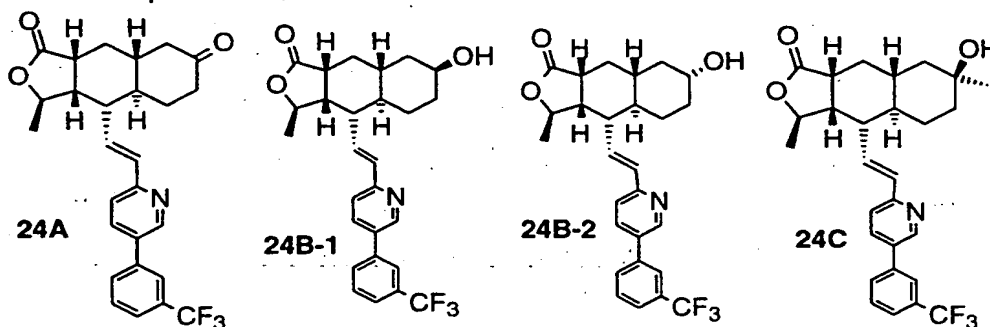
-115-

mL) and cooled to 0°C. Pd(Ph₃P)₄ (186 mg) was added, followed by tributyltinhydride (1.3 mL, 4.8 mmol). The mixture was stirred for 3 h, then chromatographed (4:1-2.5:1 hexane:EtOAc) to give 450 mg (48 %) of the desired compound. ¹H NMR (CDCl₃) δ 1.24 (d, *J* = 6.5 Hz, 3H), 1.0-1.9 (m, 10H), 2.48 (m, 1H), 2.6-2.7 (m, 2H), 3.87 (m, 4H), 4.54 (m, 1H), 9.70 (br s, 1H).

Step 8:

The product of Example 4, Step 3 (1.14 g, 3.0 mmol) was dissolved in THF (10 mL) and cooled to 0°C. *n*-BuLi (1.9 mL of 2.5 M solution in hexanes, 2.9 mmol) was added and the mixture stirred for 10 min. The solution was then added to a solution of the product of Step 7 (450 mg, 1.53 mmol) in THF (10 mL) at 0°C. The mixture was stirred for 2 h, then NH₄Cl (sat) was added. The mixture was extracted (EtOAc), dried (MgSO₄), concentrated *in vacuo* and then chromatographed (60:40 hexane:EtOAc) to give 650 mg (83 %) of the title compound. ¹H NMR (CDCl₃) δ 1.12-1.55 (m, 6H), 1.43 (d, *J* = 6 Hz, 3H), 1.78 (m, 2H), 1.79 (m, 1H), 1.96 (dd, *J* = 6.5, 3.0 Hz, 1H), 2.9 (m, 2H), 2.70 (quintet, *J* = 6.5 Hz, 1H), 3.95 (m, 4H), 4.76 (m, 1H), 6.55 (d, *J* = 15.5 Hz, 1H), 6.65 (m, 1H), 7.29 (d, *J* = 8.0 Hz, 1H), 7.60 (t, *J* = 7.5 Hz, 1H), 7.66 (d, *J* = 7.5 Hz, 1H), 7.75 (d, *J* = 7.5 Hz, 1H), 7.80 (s, 1H), 7.86 (d, *J* = 8.0 Hz, 1H), 8.79 (s, 1H).

The product of Example 24 is treated as described below to obtain Examples 24A, 24B-1, 24B-2 and 24C:



24A: The product of Example 24 (650 mg, 1.26 mmol) was dissolved in acetone (7.5 mL) and HCl (7.5 mL of a 1M solution) was added. The mixture was heated at 50°C for 16 h. NaHCO₃ (sat'd) was added and the mixture extracted with EtOAc. The combined extracts were dried

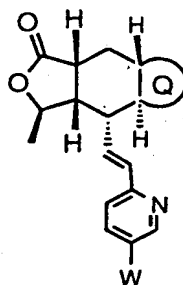
(MgSO₄), concentrated *in vacuo* and chromatographed (1:1 hexane: EtOAc) to give 590 mg (99 %) of compound 24A. ¹H NMR (CDCl₃) δ 1.2-1.5 (m, 2H), 1.47 (d, *J* = 7.0 Hz, 3H), 1.65 (m, 2H), 2.08 (m, 2H), 2.10 (m, 2H), 2.3-2.5 (m, 4H), 2.74 (quintet, *J* = 6.5 Hz, 1H), 4.80 (m, 1H),
5 6.59 (d, *J* = 6.5 Hz, 1H), 6.72 (m, 1H), 7.28 (d, *J* = 8.0 Hz, 1H), 7.61 (t, *J* = 7.5 Hz, 1H), 7.66 (d, *J* = 7.5 Hz, 1H), 7.76 (d, *J* = 7.5 Hz, 1H), 7.81 (s, 1H), 7.87 (d, *J* = 8.0 Hz, 1H), 8.80 (s, 1H).

24B-1 and 24B-2: The product of Example 24A (100 mg, 0.213 mmol) was dissolved in EtOH (8 mL) and NaBH₄ (30 mg) was added. After 5
10 min, NaHCO₃ (sat) was added and the mixture extracted with EtOAc. The extracts were dried (MgSO₄) and concentrated *in vacuo*. Purification by preparative TLC (47.5:47.5:5 hexane:EtOAc:CH₃OH) gave the least polar isomer, 24B-1 (15 mg, 15 %): ¹H NMR (CDCl₃) δ 1.15-1.4 (m, 4H), 1.43 (d, *J* = 6.0 Hz, 3H), 1.5-1.7 (m, 3H), 1.75-1.95 (m, 3H), 2.35-2.5 (m, 2H), 2.72 (quintet, *J* = 6.6 Hz, 1H), 4.16 (br s, 1H), 4.75
15 (m, 1H), 5.46, *J* = 15.5 Hz, 1H), 6.65 (m, 1H), 7.29 (d, *J* = 8.0 Hz, 1H), 7.60 (t, *J* = 8 Hz, 1H), 7.66 (d, *J* = 7.5 Hz, 1H), 7.7 δ (d, *J* = 7.5 Hz, 1H), 7.80 (s, 1H), 7.85 (d, *J* = 8.0 Hz, 1H), 8.79 (s, 1H); and the most polar isomer, 24B-2 (70 mg, 70 %): ¹H NMR (CDCl₃) δ 0.93 (m, 1H), 1.06-1.4
20 (m, 5H), 1.43 (d, *J* = 6.0 Hz, 3H), 1.85-2.05 (m, 4H), 2.40 (m, 2H), 2.70 (quintet, *J* = 6.5 Hz, 1H), 3.64 (m, 1H), 4.75 (m, 1H), 6.55 (d, *J* = 15.5 Hz, 1H), 6.64 (m, 1H), 7.29 (d, *J* = 8.0 Hz, 1H), 7.60 (t, *J* = 7.75 Hz, 1H), 7.65 (d, *J* = 7.5 Hz, 1H), 7.75 (d, *J* = 7.5 Hz, 1H), 7.80 (s, 1H), 7.85 (d, *J* = 8.0 Hz, 1H), 8.79 (s, 1H).

25 24C: The product of Example 24A (30 mg, 0.0638 mmol) was dissolved in THF (1 mL). CH₃MgBr (150 μL of a 1M solution) was added. TLC showed a more polar compound. NH₄Cl (sat'd) was added and the mixture extracted with EtOAc. The extracts were dried (MgSO₄) and concentrated *in vacuo*. The residue was purified by preparative TLC
30 (30:70 hexane:EtOAc) to give 6 mg of the desired compound. MS (FAB) *m/z* 486 (100).

Using similar procedures, compounds of the following structural formulas were prepared, wherein the variables are as defined in the table:

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Ex.	Q	W	Physical Data
24D			MS (FAB) m/z = 418
24E			MS (FAB) m/z = 418
24F			MS (FAB) m/z = 418
24G			MS (FAB) m/z = 418
24H			MS (FAB) m/z = 422
24I			MS (FAB) m/z = 422
24J			MS (FAB) m/z = 422
24K			MS (FAB) m/z = 422
24L			^1H NMR (CDCl_3) 0.93 (m, 1H), 1.05-1.38 (m, 5H), 1.43 (d, J = 6.0 Hz, 3H), 1.88-2.1 (m, 4H), 2.36 (m, 2H), 2.68 (m, 1H), 3.65 (m, 1H), 5.75 (m, 1H), 6.5-6.65 (m, 2H), 7.27 (d, J = 8.0 Hz, 1H), 7.35-7.5 (m, 3H), 7.55 (s, 1H), 7.81 (d, J = 8.0 Hz, 1H), 8.75 (s, 1H).

The following formulations exemplify some of the dosage forms of this invention. In each, the term "active compound" designates a compound of formula I.

5

EXAMPLE A - Tablets

<u>No.</u>	<u>Ingredient</u>	<u>mg/tablet</u>	<u>mg/tablet</u>
1	Active Compound	100	500
2	Lactose USP	122	113
3	Corn Starch, Food Grade, as a 10% paste in Purified Water	30	40
4	Corn Starch, Food Grade	45	40
5	Magnesium Stearate	<u>3</u>	<u>7</u>
Total		300	700

Method of Manufacture

- Mix Item Nos. 1 and 2 in suitable mixer for 10-15 minutes. Granulate the mixture with Item No. 3. Mill the damp granules through a coarse screen (e.g., 1/4", 0.63 cm) if necessary. Dry the damp granules. Screen the dried granules if necessary and mix with Item No. 4 and mix for 10-15 minutes. Add Item No. 5 and mix for 1-3 minutes. Compress the mixture to appropriate size and weight on a suitable tablet machine.

10

EXAMPLE B - Capsules

<u>No.</u>	<u>Ingredient</u>	<u>mg/tablet</u>	<u>mg/tablet</u>
1	Active Compound	100	500
2	Lactose USP	106	123
3	Corn Starch, Food Grade	40	70
4	Magnesium Stearate NF	<u>4</u>	<u>7</u>
Total		250	700

15 Method of Manufacture

Mix Item Nos. 1, 2 and 3 in a suitable blender for 10-15 minutes. Add Item No. 4 and mix for 1-3 minutes. Fill the mixture into suitable two-piece hard gelatin capsules on a suitable encapsulating machine.

20

The activity of the compounds of formula I can be determined by the following procedures.

In Vitro Testing Procedure for Thrombin Receptor Antagonists:Preparation of [³H]haTRAP

A(pF-F)R(ChA)(hR)(I₂-Y)-NH₂ (1.03 mg) and 10% Pd/C (5.07 mg) were suspended in DMF (250 µl) and diisopropylethylamine (10 µl).

- 5 The vessel was attached to the tritium line, frozen in liquid nitrogen and evacuated. Tritium gas (342 mCi) was then added to the flask, which was stirred at room temperature for 2 hours. At the completion of the reaction, the excess tritium was removed and the reacted peptide solution was diluted with DMF (0.5 ml) and filtered to remove the catalyst. The collected DMF solution of the crude peptide was diluted with water and freeze dried to remove the labile tritium. The solid peptide was redissolved in water and the freeze drying process repeated. The tritiated peptide ([³H]haTRAP) was dissolved in 0.5 ml of 0.1% aqueous TFA and purified by HPLC using the following conditions:
- 10 column, Vydac C18, 25 cm x 9.4 mm I.D.; mobile phase, (A) 0.1% TFA in water, (B) 0.1% TFA in CH₃CN; gradient, (A/B) from 100/0 to 40/60 over 30 min; flow rate, 5 ml /min; detection, UV at 215 nm. The radiochemical purity of [³H]haTRAP was 99% as analyzed by HPLC. A batch of 14.9 mCi at a specific activity of 18.4 Ci/mmol was obtained.

20 Preparation of platelet membranes

- Platelet membranes were prepared using a modification of the method of Natarajan et al (Natarajan et al, Int. J. Peptide Protein Res. 45:145-151 (1995)) from 20 units of platelet concentrates obtained from the North Jersey Blood Center (East Orange, NJ) within 48 hours of collection. All steps were carried out at 4° C under approved biohazard safety conditions. Platelets were centrifuged at 100 x g for 20 minutes at 4° C to remove red cells. The supernatants were decanted and centrifuged at 3000 x g for 15 minutes to pellet platelets. Platelets were resuspended in 10 mM Tris-HCl, pH 7.5, 150 mM NaCl, 5 mM EDTA, to a total volume of 200 ml and centrifuged at 4400 x g for 10 minutes. This step was repeated two additional times. Platelets were resuspended in 5 mM Tris-HCl, pH 7.5, 5 mM EDTA to a final volume of approximately 30 ml and were homogenized with 20 strokes in a Dounce homogenizer. Membranes were pelleted at 41,000 x g, resuspended in 40-50 ml 20 mM Tris-HCl, pH 7.5, 1 mM EDTA, 0.1 mM dithiothreitol, and 10 ml aliquots were frozen in liquid N₂ and stored at -80° C. To complete

membrane preparation, aliquots were thawed, pooled, and homogenized with 5 strokes of a Dounce homogenizer. Membranes were pelleted and washed 3 times in 10 mM triethanolamine-HCl, pH 7.4, 5 mM EDTA, and resuspended in 20-25 ml 50 mM Tris-HCl, pH 7.5, 10 mM MgCl₂, 1 mM EGTA, and 1% DMSO. Aliquots of membranes were frozen in liquid N₂ and stored at -80° C. Membranes were stable for at least 3 months. 20 units of platelet concentrates typically yielded 250 mg of membrane protein. Protein concentration was determined by a Lowry assay (Lowry et al, J. Biol. Chem., 193:265-275 (1951)).

10 High Throughput Thrombin Receptor Radioligand Binding Assay

Thrombin receptor antagonists were screened using a modification of the thrombin receptor radioligand binding assay of Ahn et al. (Ahn et al, Mol. Pharmacol., 51:350-356 (1997)). The assay was performed in 96 well Nunc plates (Cat. No. 269620) at a final assay volume of 200 µl. Platelet membranes and [³H]haTRAP were diluted to 0.4 mg/ml and 22.2 nM, respectively, in binding buffer (50 mM Tris-HCl, pH 7.5, 10 mM MgCl₂, 1 mM EGTA, 0.1% BSA). Stock solutions (10 mM in 100% DMSO) of test compounds were further diluted in 100% DMSO. Unless otherwise indicated, 10 µl of diluted compound solutions and 90 µl of radioligand (a final concentration of 10 nM in 5% DMSO) were added to each well, and the reaction was started by the addition of 100 µl of membranes (40 µg protein/well). The binding was not significantly inhibited by 5% DMSO. Compounds were tested at three concentrations (0.1, 1 and 10 µM). The plates were covered and vortex-mixed gently on a Lab-Line Titer Plate Shaker for 1 hour at room temperature. Packard UniFilter GF/C filter plates were soaked for at least 1 hour in 0.1% polyethyleneimine. The incubated membranes were harvested using a Packard FilterMate Universal Harvester and were rapidly washed four times with 300 µl ice cold 50 mM Tris-HCl, pH 7.5, 10 mM MgCl₂, 1 mM EGTA. MicroScint 20 scintillation cocktail (25 µl) was added to each well, and the plates were counted in a Packard TopCount Microplate Scintillation Counter. The specific binding was defined as the total binding minus the nonspecific binding observed in the presence of excess (50 µM) unlabeled haTRAP. The % inhibition by a compound of [³H]haTRAP binding to thrombin receptors was calculated from the following relationship:

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% Inhibition =

$$\frac{\text{Total binding} - \text{Binding in the presence of a test compound}}{\text{Total binding} - \text{Nonspecific binding}} \times 100$$

Materials

5 A(pF-F)R(ChA)(hR)Y-NH₂ and A(pF-F)R(ChA)(hR)(I₂-Y)-NH₂, were custom synthesized by AnaSpec Inc. (San Jose, CA). The purity of these peptides was >95%. Tritium gas (97%) was purchased from EG&G Mound, Miamisburg Ohio. The gas was subsequently loaded and stored on an IN/US Systems Inc. Trisorber. MicroScint 20
10 scintillation cocktail was obtained from Packard Instrument Co.

Protocol For Ex-Vivo Platelet Aggregation In Cynomolgus Whole Blood Drug administration and blood collection:

 Conscious chaired cynomolgus monkeys are allowed to equilibrate for 30 min. A needle catheter is inserted into a brachial vein
15 for infusion of test drugs. Another needle catheter is inserted into the other brachial or saphenous vein and used for blood sampling. In those experiments where the compound is administered orally only one catheter is used. A baseline blood sample (1-2 ml) is collected in vacutainer tubes containing a thrombin inhibitor CVS 2139 (100 µg/0.1
20 ml saline) as an anticoagulant. The drug is then infused intravenously over a period of 30 min. Blood samples (1 ml) are collected at 5, 10, 20, 30 min during and 30, 60, 90 min after termination of the drug infusion. In PO experiments the animals are dosed with the drug using a gavage cannula. Blood samples are collected at 0, 30, 60, 90, 120, 180, 240,
25 300, 360 min after dosing. 0.5 ml of the blood is used for whole blood aggregation and the other 0.5 ml is used for determining the plasma concentration of the drug or its metabolites. Aggregation is performed immediately after collection of the blood sample as described below.

Whole Blood Aggregation:

30 A 0.5 ml blood sample is added to 0.5 ml of saline and warmed to 37°C in a Chronolog whole blood aggregometer. Simultaneously, the impedance electrode is warmed in saline to 37°C. The blood sample with a stir bar is placed in the heating block well, the impedance electrode is placed in the blood sample and the collection software is started. The
35 software is allowed to run until the baseline is stabilized and then a 20 Ω calibration check is performed. 20 Ω is equal to 4 blocks on the graphic

produced by the computer software. The agonist (haTRAP) is added by an adjustable volume pipette (5-25 μ l) and the aggregation curve is recorded for 10 minutes. Maximum aggregation in 6 minutes following agonist is the value recorded.

5 *In vitro* Platelet Aggregation Procedure:

Platelet aggregation studies were performed according to the method of Bednar *et al.* (Bednar, B., Condra, C., Gould, R.J., and Connolly, T.M., Throm. Res., 77:453-463 (1995)). Blood was obtained from healthy human subjects who were aspirin free for at least 7 days by
10 venipuncture using ACD as anticoagulant. Platelet rich plasma was prepared by centrifugation at 100Xg for 15 minutes at 15 deg C. Platelets were pelleted at 3000xg and washed twice in buffered saline containing 1 mM EGTA and 20 μ g/ml apyrase to inhibit aggregation. Aggregation was performed at room temperature in buffered saline
15 supplemented with 0.2 mg/ml human fibrinogen. Test compound and platelets were preincubated in 96-well flat-bottom plates for 60 minutes. Aggregation was initiated by adding 0.3 μ M haTRAP or 0.1 U/ml thrombin and rapidly vortexing the mixture using a Lab Line Titer Plate Shaker (speed 7). Percent aggregation was monitored as increasing
20 light transmittance at 405 nm in a Spectromax Plate Reader.

In vivo Antitumor Procedure:

Tests in the human breast carcinoma model in nude mouse are conducted according to the procedure reported in S. Even-Ram *et. al.*, Nature Medicine, 4, 8 (1998), p. 909-914.

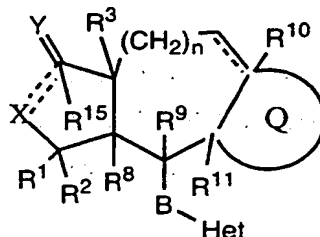
25 Using the test procedures described above, in the *in vitro* thrombin receptor antagonist assay, compounds of the invention were found to have IC₅₀ values (i.e., the concentration at which a 50% inhibition of thrombin receptor was observed) in the range of about 4 to 2000 nM, with preferred compounds having IC₅₀ values in the range of
30 about 4 to 100 nM.

In the *in vitro* Platelet Aggregation Inhibition (PAI) assay, compounds tested were found to have IC₅₀ values in the range of 67 to 1000 nM. In the *ex-vivo* C. Monkey Whole Blood PAI, one compound tested showed 100% aggregation at 3 mpk (oral in betahydroxypropyl-
35 cyclodextrin as cosolvent) and another showed 100% aggregation at 10 mpk (i.v. infusion over 30 min. in 5% dextrose).

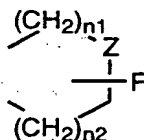
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What is claimed:

1. A compound represented by the structural formula



- 5 or a pharmaceutically acceptable salt thereof, wherein:
 the single dotted line represents an optional double bond;
 the double dotted line represents an optional single bond;
 n is 0-2;
 Q is



10

wherein n_1 and n_2 are independently 0-2; or when the double bond is not present, Q is also fused R-substituted aryl or R-substituted heteroaryl;

- 15 R is 1 to 3 substituents independently selected from the group consisting of H, C₁-C₆ alkyl, halogen, hydroxy, amino, (C₁-C₆)alkyl-amino, (C₁-C₆)dialkylamino, (C₁-C₆)alkoxy, -COR¹⁶, -COOR¹⁷, -SOR¹⁶, -SO₂R¹⁶, -NR¹⁶COR^{16a}, -NR¹⁶COOR^{16a}, -NR¹⁶CONR⁴R⁵, fluoro-(C₁-C₆)alkyl, difluoro(C₁-C₆)alkyl, trifluoro(C₁-C₆)alkyl, C₃-C₆ cycloalkyl, C₂-C₆ alkenyl, aryl(C₁-C₆)alkyl, aryl(C₂-C₆)alkenyl, heteroaryl(C₁-C₆)-alkyl, heteroaryl(C₂-C₆)alkenyl, hydroxy(C₁-C₆)alkyl, amino(C₁-C₆)-alkyl, aryl and thio(C₁-C₆)alkyl;

- 20 R¹ and R² are independently selected from the group consisting of H, C₁-C₆ alkyl, fluoro(C₁-C₆)alkyl, difluoro(C₁-C₆)alkyl, trifluoro-(C₁-C₆)alkyl, C₃-C₆ cycloalkyl, C₂-C₆ alkenyl, aryl(C₁-C₆)alkyl, aryl(C₂-C₆)alkenyl, heteroaryl(C₁-C₆)alkyl, heteroaryl(C₂-C₆)alkenyl, hydroxy-(C₁-C₆)alkyl, amino(C₁-C₆)alkyl, aryl and thio(C₁-C₆)alkyl; or R¹ and R² together form a =O group;

R³ is H, hydroxy, C₁-C₆ alkoxy, -SOR¹⁶, -SO₂R¹⁷, -C(O)OR¹⁷, -C(O)NR¹⁸R¹⁹, C₁-C₆ alkyl, halogen, fluoro(C₁-C₆)alkyl, difluoro(C₁-

C₆)alkyl, trifluoro(C₁-C₆)alkyl, C₃-C₆ cycloalkyl, C₂-C₆ alkenyl, aryl(C₁-C₆)alkyl, aryl(C₂-C₆)alkenyl, heteroaryl(C₁-C₆)alkyl, heteroaryl(C₂-C₆)alkenyl, hydroxy(C₁-C₆)alkyl, amino(C₁-C₆)alkyl, aryl or thio(C₁-C₆)alkyl;

- 5 Het is a mono-, bi- or tricyclic heteroaromatic group of 5 to 14 atoms comprised of 1 to 13 carbon atoms and 1 to 4 heteroatoms independently selected from the group consisting of N, O and S, wherein a ring nitrogen can form an N-oxide or a quaternary group with a C₁-C₄ alkyl group, wherein Het is attached to B by a carbon atom ring member, and wherein the Het group is substituted by 1 to 4 substituents, W, independently selected from the group consisting of H; C₁-C₆ alkyl; fluoro(C₁-C₆)alkyl; difluoro(C₁-C₆)alkyl; trifluoro-(C₁-C₆)-alkyl; C₃-C₆ cycloalkyl; heterocycloalkyl; heterocycloalkyl substituted by C₁-C₆ alkyl or C₂-C₆ alkenyl; C₂-C₆ alkenyl; R²¹-aryl(C₁-C₆)alkyl; R²¹-aryl-(C₂-C₆)-alkenyl; heteroaryl(C₁-C₆)alkyl; heteroaryl(C₂-C₆)-alkenyl; hydroxy(C₁-C₆)alkyl; dihydroxy(C₁-C₆)alkyl; amino(C₁-C₆)alkyl; (C₁-C₆)alkylamino-(C₁-C₆)alkyl; di-((C₁-C₆)alkyl)-amino(C₁-C₆)alkyl; thio(C₁-C₆)alkyl; C₁-C₆ alkoxy; C₂-C₆ alkenyloxy; halogen; -NR⁴R⁵; -CN; -OH; -COOR¹⁷; -COR¹⁶; -OSO₂CF₃; -CH₂OCH₂CF₃; (C₁-C₆)alkylthio; -C(O)NR⁴R⁵; -OCHR⁶-phenyl; phenoxy-(C₁-C₆)alkyl; -NHCOR¹⁶; -NH₂SO₂R¹⁶; biphenyl; -OC(R⁶)₂COOR⁷; -OC(R⁶)₂C(O)NR⁴R⁵; (C₁-C₆)alkoxy; C₁-C₆ alkoxy substituted by (C₁-C₆)alkyl, amino, -OH, COOR¹⁷, -NHCOOR¹⁷, -CONR⁴R⁵, aryl, aryl substituted by 1 to 3 substituents independently selected from the group consisting of halogen, -CF₃, C₁-C₆ alkyl, C₁-C₆ alkoxy and -COOR¹⁷, aryl wherein adjacent carbons form a ring with a methylenedioxy group, -C(O)NR⁴R⁵ and heteroaryl; R²¹-aryl; aryl wherein adjacent carbons form a ring with a methylenedioxy group; heteroaryl; heteroaryl substituted by 1 to 4 substituents selected from the group consisting of halogen, C₁-C₆ alkoxy, C₁-C₆ alkyl, (C₁-C₆)alkyl-amino, di-((C₁-C₆)alkyl)amino, -OCF₃, -NO₂, hydroxy(C₁-C₆)alkyl, -CHO and phenyl; and heteroaryl wherein adjacent carbon atoms form a ring with a C₃-C₅ alkylene group or a methylenedioxy group;
- 35 R⁴ and R⁵ are independently selected from the group consisting of H, C₁-C₆ alkyl, phenyl, benzyl and C₃-C₆ cycloalkyl, or R⁴ and R⁵

together are $-(CH_2)_4-$, $-(CH_2)_5-$ or $-(CH_2)_2NR^7-(CH_2)_2-$ and form a ring with the nitrogen to which they are attached;


R^6 is independently selected from the group consisting of H, C_1 - C_6 alkyl or phenyl;

5 R^7 is H or (C_1-C_6) alkyl;

R^8 , R^{10} and R^{11} are independently selected from the group consisting of R^1 and $-OR^1$, provided that when the optional double bond is present, R^{10} is absent, and when ring Q is aromatic, R^{10} and R^{11} are absent;

10 R^9 is H, OH, C_1 - C_6 alkoxy, halogen or halo(C_1 - C_6)alkyl;

B is $-(CH_2)_{n3}-$, $-CH_2-O-$, $-CH_2S-$, $-CH_2-NR^6-$, $-C(O)NR^6-$.

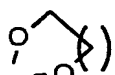
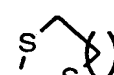
$-NR^6C(O)-$, , cis or trans $-(CH_2)_{n4}CR^{12}=CR^{12a}(CH_2)_{n5}$ or $-(CH_2)_{n4}C\equiv C(CH_2)_{n5}-$, wherein n_3 is 0-5, n_4 and n_5 are independently 0-2, and R^{12} and R^{12a} are independently selected from the group

15 consisting of H, C_1 - C_6 alkyl and halogen;

X is $-O-$ or $-NR^6-$ when the double dotted line represents a single bond, or X is $-OH$ or $-NHR^{20}$ when the bond is absent;

20 Y is $=O$, $=S$, (H, H), (H, OH) or (H, C_1 - C_6 alkoxy) when the double dotted line represents a single bond, or when the bond is absent, Y is $=O$, (H, H), (H, OH), (H, SH) or (H, C_1 - C_6 alkoxy);

R^{15} is absent when the double dotted line represents a single bond and is H, C_1 - C_6 alkyl, $-NR^{18}R^{19}$, or $-OR^{17}$ when the bond is

absent; or Y is  $_{1-2}$ or  $_{1-2}$ and R^{15} is H or C_1 - C_6 alkyl;

25 R^{16} and R^{16a} are independently selected from the group consisting of C_1 - C_6 lower alkyl, phenyl or benzyl;

R^{17} , R^{18} and R^{19} are independently selected from the group consisting of H, C_1 - C_6 alkyl, phenyl, benzyl;

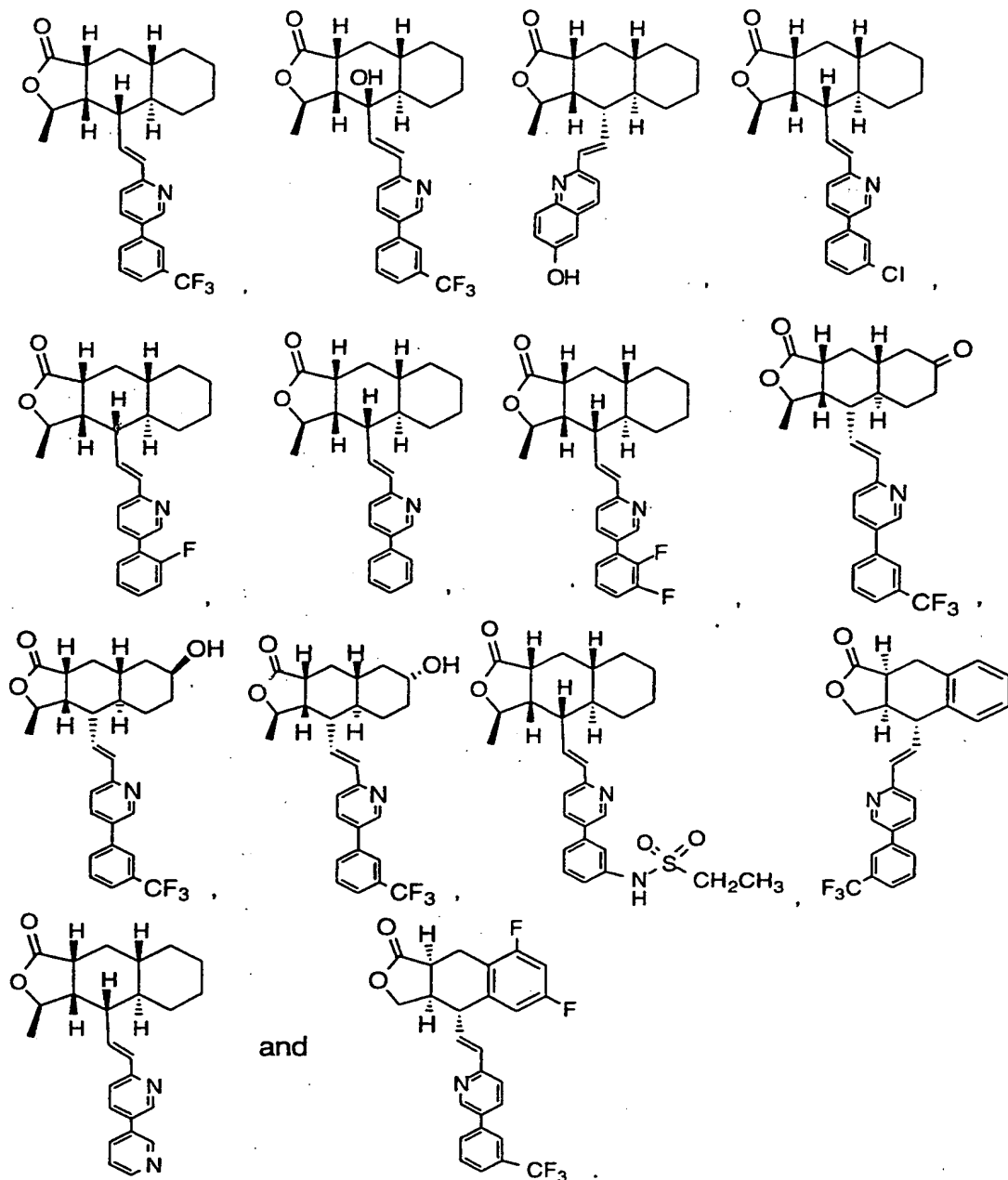
R^{20} is H, C_1 - C_6 alkyl, phenyl, benzyl, $-C(O)R^6$ or $-SO_2R^6$;

30 R^{21} is 1 to 3 substituents independently selected from the group consisting of $-CF_3$, $-OCF_3$, halogen, $-NO_2$, C_1 - C_6 alkyl, C_1 - C_6 alkoxy, (C_1-C_6) alkylamino, di- $((C_1-C_6)$ alkyl)amino, amino(C_1 - C_6)alkyl, (C_1-C_6) -alkylamino(C_1 - C_6)alkyl, di- $((C_1-C_6)$ alkyl)-amino(C_1 - C_6)alkyl, hydroxy- (C_1-C_6) alkyl, $-COOR^{17}$, $-COR^{17}$, $-NHCOR^{16}$, $-NHSO_2R^{16}$ and $-NHSO_2CH_2CF_3$;

Z is -CH₂-, -O-, -S(O)₀₋₂-, -NR²²-, -C(O)-, -C(=NOR¹⁷)- or -C(R¹³R¹⁴)-, wherein R¹³ and R¹⁴, together with the carbon to which they are attached, form a spirocycloalkyl group of 3 to 6 carbons, or a spiroheterocycloalkyl group of 3 to 6 members, comprised of 2 to 5 carbon atoms and 1 or 2 heteroatoms selected from the group consisting of O, S and N; and
R²² is H, C₁-C₆ alkyl, phenyl, benzyl, -COR¹⁶ or -CONR¹⁸R¹⁹.

2. A compound of claim 1 wherein n is zero.
3. A compound of any of claims 1 or 2 wherein the double dotted line represents a single bond, X is -O- and Y is =O.
4. A compound of any of claims 1, 2 or 3 wherein R² and R⁸ are each hydrogen; R³ is hydrogen or C₁-C₆ alkyl; and R⁹ is H, OH or C₁-C₆ alkoxy.
5. A compound of any of claims 1, 2, 3 or 4 wherein R¹ is C₁-C₆ alkyl.
6. A compound of any of claims 1, 2, 3, 4 or 5 wherein Q is R-substituted cyclohexyl or R-substituted phenyl.
7. A compound of any of claims 1, 2, 3, 4, 5 or 6 wherein R is H, fluorine, OH, C₁-C₆ alkyl, C₁-C₆ alkoxy or -COOR¹⁷.
8. A compound of any of claims 1, 2, 3, 4, 5, 6 or 7 wherein B is -CH=CH-.
9. A compound of any of claims 1, 2, 3, 4, 5, 6, 7 or 8 wherein Het is pyridyl, W-substituted pyridyl, quinolyl or W-substituted quinolyl.
10. A compound of claim 9 wherein W is C₁-C₆ alkyl, aryl, R²¹-aryl or heteroaryl.
11. A compound of claim 1 selected from the group consisting of

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5

12. A pharmaceutical composition comprising an effective amount of a compound of claim 1 and a pharmaceutically acceptable carrier.

10 13. A method of inhibiting thrombin receptors comprising administering to a mammal in need of such treatment an effective amount of a compound of claim 1.

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14. A method of treating thrombosis, atherosclerosis, restenosis, hypertension, angina pectoris, arrhythmia, heart failure, myocardial infarction, glomerulonephritis, thrombotic stroke, thromboembolytic stroke, peripheral vascular diseases, inflammatory disorders, cerebral ischemia and cancer, comprising administering to a mammal in need of such treatment an effective amount of a compound of claim 1.
- 5

INTERNATIONAL SEARCH REPORT

International Application No
PCT/US 98/24523

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 C07D405/06 C07D405/14 C07D409/14 C07D405/12 C07D417/06
C07D401/06 C07D491/04 C07D471/04 C07D493/10 A61K31/44
A61K31/47 //(C07D491/04,317:00,221:00),(C07D471/04,221:00,

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C07D A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	S. CHACKALAMANNIL ET AL.: "A highly efficient total synthesis of (+)-Himbacine" JOURNAL OF THE AMERICAN CHEMICAL SOCIETY., vol. 118, no. 40, 9 October 1996, pages 9812-9813, XP002095052 DC US cited in the application see the whole document ----- -/--	1-10

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents :

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

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- "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- "&" document member of the same patent family

Date of the actual completion of the international search

1 March 1999

Date of mailing of the international search report

18/03/1999

Name and mailing address of the ISA

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INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 98/24523

A. CLASSIFICATION OF SUBJECT MATTER
IPC 6 221:00), (C07D493/10, 317:00, 307:00)

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevance to claim No
A	M.S. BERNATOWICZ ET AL.: "Development of potent thrombin receptor antagonist peptides" JOURNAL OF MEDICINAL CHEMISTRY., vol. 39, no. 25, 6 December 1996, pages 4879-4887, XP002095053 WASHINGTON US cited in the application see the whole document	1,12
A	WO 94 03479 A (COR THERAPEUTICS, INC.) 17 February 1994 cited in the application see claims	1,12

☐ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents:

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- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

- "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
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- "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- "G" document member of the same patent family

Date of the actual completion of the international search

1 March 1999

Date of mailing of the international search report

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Authorized officer

Chouly, J

INTERNATIONAL SEARCH REPORT

international application No.

PCT/US 98/ 24523

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.: 13,14
because they relate to subject matter not required to be searched by this Authority, namely:
Remark: Although claims 13.14
are directed to a method of treatment of the human/animal
body, the search has been carried out and based on the alleged
effects of the compound/composition.
2. ☐ Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such
an extent that no meaningful international Search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all
searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment
of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report
covers only those claims for which fees were paid, specifically claims Nos
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is
restricted to the invention first mentioned in the claims; it is covered by claims Nos

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No.

PCT/US 98/24523

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9403479 A	17-02-1994	AU 669922 B	27-06-1996
		AU 4668593 A	03-03-1994
		CA 2140543 A	17-02-1994
		EP 0656008 A	07-06-1995
		JP 2675438 B	12-11-1997
		JP 7506115 T	06-07-1995
		SG 49772 A	15-06-1998
		US 5866681 A	02-02-1999

Form PCT/ISA/210 (patent family annex) (July 1992)